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(88) Date of publication of the international search report:

18 April 2002

INTERNATIONAL SEARCH REPORT

national Application No
PCT/US 01/09341

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C12N15/12 C07K14/435 C07K14/705 C07K16/18 C12N5/10
C12Q1/68 G01N33/50

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07K G01N C12N C12Q

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

SEQUENCE SEARCH, EPO-Internal, BIOSIS, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE EMBL SEQUENCE LIBRARY [Online] 6 January 2000 (2000-01-06) ADAMS, M. AND VENTER, J.C.: "Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in ordered pieces" XP002182628 accession no AC020076	1,2,4,5
A	--- WO 94 08006 A (ZYMOGENETICS INC) 14 April 1994 (1994-04-14) the whole document --- -/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

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"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

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INTERNATIONAL SEARCH REPORT

national Application No

PCT/US 01/09341

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>LI X-J ET AL: "CLONING, FUNCTIONAL EXPRESSION, AND DEVELOPMENTAL REGULATION OF A NEUROPEPTIDE Y RECEPTOR FROM DROSOPHILA MELANOGASTER" JOURNAL OF BIOLOGICAL CHEMISTRY, AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD, US, vol. 267, no. 1, 5 January 1992 (1992-01-05), pages 9-12, XP000877443 ISSN: 0021-9258 the whole document</p> <p>---</p>	
A	<p>WO 99 01468 A (DEN HEUVEL MARCEL VAN ;INGHAM PHILIP W (GB); ONTOGENY INC (US)) 14 January 1999 (1999-01-14) the whole document</p> <p>---</p>	
A	<p>HAUSER FRANK ET AL: "Molecular cloning, genomic organization and developmental regulation of a novel receptor from Drosophila melanogaster structurally related to gonadotropin-releasing hormone receptors from vertebrates." BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, vol. 249, no. 3, 28 August 1998 (1998-08-28), pages 822-828, XP002182627 ISSN: 0006-291X the whole document</p> <p>---</p>	
A	<p>FENG G ET AL: "CLONING AND FUNCTIONAL CHARACTERIZATION OF A NOVEL DOPAMINE RECEPTOR FROM DROSOPHILA MELANOGASTER" JOURNAL OF NEUROSCIENCE, NEW YORK, NY, US, vol. 15, no. 12, 15 June 1995 (1995-06-15), pages 3925-3933, XP002919142 ISSN: 0270-6474 the whole document</p> <p>---</p>	
P,X	<p>ADAMS M D ET AL: "THE GENOME SEQUENCE OF DROSOPHILA MELANOGASTER" SCIENCE, AMERICAN ASSOCIATION FOR THE ADVANCEMENT OF SCIENCE,, US, vol. 287, no. 5461, 24 March 2000 (2000-03-24), pages 2185-2195, XP000961051 ISSN: 0036-8075 the whole document</p> <p>---</p> <p style="text-align: center;">-/--</p>	1,2,4,5

INTERNATIONAL SEARCH REPORT

national Application No
PCT/US 01/09341

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	<p>DATABASE EMBL SEQUENCE LIBRARY [Online] 24 March 2000 (2000-03-24) ADAMS, M.D.; ETAL.: "Drosophila melanogaster genomic scaffold 142000013386053 section 8 of 30 complete sequence" XP002182629 accession no. AE003491 and AE002593</p> <p>---</p>	1,2,4,5
P,X	<p>DATABASE TREMBLREL. DATABASE [Online] 1 May 2000 (2000-05-01) ADAMS, M.D., ET AL.: "The genome sequence of Drosophila melanogaster - cross-reference to EMBL accession no. AE003491, AAF48216.1" XP002182630 accession no. Q9VYH9</p> <p>---</p>	1,2
E	<p>EP 1 136 501 A (BAYER AG) 26 September 2001 (2001-09-26) the whole document</p> <p>-----</p>	1-20

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 01/09341

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-20 partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

Invention 1: claims 1-20 partially

Isolated protein consisting or comprising an amino acid sequence as characterized by SEQID3, or an allelic variant or an ortholog of said amino acid sequence wherein said variant or ortholog is encoded by a nucleic acid molecule that hybridizes to the nucleic acid molecule as characterized by SEQID1 or 2; an antibody that binds to said protein; furthermore a nucleic acid molecule consisting or comprising of a nucleotide sequence that

- 1) encodes the amino acid sequence of SEQID3
- 2) encodes and allelic variant or an ortholog of an amino acid sequence of SEQID3 wherein said nucleotide sequence hybridizes to SEQID 1 or 2
- 3) encodes a fragment of said SEQID3
- 4) is the complement of the nucleotides of 1) to 3)

The recombinant expression of the same in host cells and methods for the detection of said proteins or said nucleic acids in a sample with the help of an agent that binds to said protein or an oligonucleotide and kits that contain such agent or oligonucleotide.

Furthermore, a method to identify an agent that binds to said protein by detecting a complex formed by an agent and the said protein.

Invention 2-66: claims 1-20 partially

as invention one but referring to the protein and nucleic acid sequences as characterized by SEQIDs 6,9,....,192,195,198; SEQIDs 4,7,....,190,193,196 and SEQIDs 5,8,....,191,194,197, respectively.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 01/09341

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9408006	A	14-04-1994	CA 2145866 A1	14-04-1994
			EP 0663006 A1	19-07-1995
			JP 8501942 T	05-03-1996
			WO 9408006 A1	14-04-1994
			US 5683884 A	04-11-1997
			US 5674981 A	07-10-1997
			US 5622839 A	22-04-1997
			US 5674689 A	07-10-1997

WO 9901468	A	14-01-1999	AU 8380898 A	25-01-1999
			WO 9901468 A2	14-01-1999

EP 1136501	A	26-09-2001	DE 10013618 A1	20-09-2001
			EP 1136501 A2	26-09-2001
			JP 2001299369 A	30-10-2001

CORRECTED VERSION

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International Bureau**



(43) International Publication Date
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G01N 33/50

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(74) Agent: MILLMAN, Robert, A.; Celcra Genomics Corp., 45 West Gude Drive C2-4, Rockville, MD 20850 (US).

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(26) Publication Language: English

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09/618,893	18 July 2000 (18.07.2000)	US

[Continued on next page]

(54) Title: ISOLATED G-PROTEIN COUPLED RECEPTORS, NUCLEIC ACID MOLECULES ENCODING GPCR PROTEINS, AND USES THEREOF AS INSECTICIDAL TARGETS

Colera Sequence No. : 142000012783463

[illegible]

Exon: 4236..4062
Exon: 3530..3230
Exon: 2674..2584
Exon: 2518..2427
Exon: 2325..1679
Exon: 1380..1001
Start ATG: 4216 (Reverse strand: CAT)

TRANSCRIPT No. : CT4121

[illegible]

(57) Abstract: The present invention provides amino acid sequences of proteins that are encoded by genes within the *Drosophila melanogaster* genome, the GPCR proteins of the present invention. The present invention specifically provides isolated protein and nucleic acid molecules, methods of identifying orthologs and paralogs of the GPCR proteins and methods of identifying modulators of the GPCR proteins for use as insecticides.

WO 01/070980 A3



(84) **Designated States (regional):** ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

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**ISOLATED G-PROTEIN COUPLED RECEPTORS, NUCLEIC ACID MOLECULES
ENCODING GPCR PROTEINS, AND USES THEREOF AS INSECTICIDAL TARGETS**

RELATED APPLICATIONS

5 The present application claims priority to U.S. Serial No. 60/191,638, filed March 23, 2000 (Atty. Docket CL000388) and U.S. Serial No. 09/618,893, filed July 18, 2000 (Atty. Docket CL000733).

FIELD OF THE INVENTION

10 The present invention is in the field of G-Protein coupled receptors (GPCRs), recombinant DNA molecules and protein production. The present invention specifically provides novel GPCR proteins and nucleic acid molecules encoding such protein molecules, for use in the development of insecticide and insecticide targets and as a source for identifying human therapeutics and human therapeutic development.

15 **BACKGROUND OF THE INVENTION**

 The *Drosophila melanogaster* genome is 165 Mb, with about 120 Mb of this being euchromatic. The genome is organized in 4 chromosome pairs and is estimated to contain 10 - 12,000 genes. Model organisms, such as *Drosophila melanogaster*, share many genes with
20 humans whose sequences and functions have been conserved. In addition to myriad similarities in cellular structure and function, humans and *Drosophila* share pathways for intercellular signaling, developmental patterning, learning and behavior, as well as tumor formation and metastasis.

 The genes involved in the development of *Drosophila*, with few exceptions, are the same
25 as those involved in the development of higher organisms. Developmental biology studies the sequential activation and interaction of genes, in relation to developing morphology. Right now, *Drosophila* is the only organism for which one can begin with a list of genes active in the egg and follow the morphological changes and gene activations through to adulthood.

Drosophila studies have provided the widest knowledge base available for any single
30 organism; accordingly, developmental biologists use the fly to ferret out the activity of genes with similar functions in higher organisms. Despite its small size, the fly is by no means a small developmental problem. If you know the genes involved in the development of the fly, you also

know, to a reasonable approximation, the genes involved in the development of the worm, the fish, the mouse, and humans.

A major goal in drug/insecticide development is to understand and elucidate the molecular mechanisms that govern cell signaling and cell-cell interactions in higher eukaryotes. GPCRs form major links in cellular communication/response systems. A complete list of GPCR proteins from *Drosophila* would therefore be invaluable in developing human therapeutic targets. Not only will the proteins serve as models for human cellular signaling and response, such molecules will also serve as molecular keys in identifying therapeutically important human orthologs.

G-protein coupled receptors

G-protein coupled receptors (GPCRs) constitute a major class of proteins responsible for transducing a signal within a cell. GPCRs have three structural domains: an amino terminal extracellular domain, a transmembrane domain containing seven transmembrane segments, three extracellular loops, and three intracellular loops, and a carboxy terminal intracellular domain. Upon binding of a ligand to an extracellular portion of a GPCR, a signal is transduced within the cell that results in a change in a biological or physiological property of the cell. GPCRs, along with G-proteins and effectors (intracellular enzymes and channels modulated by G-proteins), are the components of a modular signaling system that connects the state of intracellular second messengers to extracellular inputs.

GPCR genes and gene-products are potential causative agents of disease (Spiegel *et al.*, *J. Clin. Invest.* 92:1119-1125 (1993); McKusick *et al.*, *J. Med. Genet.* 30:1-26 (1993)). Specific defects in the rhodopsin gene and the V2 vasopressin receptor gene have been shown to cause various forms of retinitis pigmentosa (Nathans *et al.*, *Annu. Rev. Genet.* 26:403-424(1992)), and nephrogenic diabetes insipidus (Holtzman *et al.*, *Hum. Mol. Genet.* 2:1201-1204 (1993)). These receptors are of critical importance to both the central nervous system and peripheral physiological processes. Evolutionary analyses suggest that the ancestor of these proteins originally developed in concert with complex body plans and nervous systems.

The GPCR protein superfamily can be divided into five families: Family I, receptors typified by rhodopsin and the β 2-adrenergic receptor and currently represented by over 200 unique members (Dohlman *et al.*, *Annu. Rev. Biochem.* 60:653-688 (1991)); Family II, the parathyroid hormone/calcitonin/secretin receptor family (Juppner *et al.*, *Science* 254:1024-1026 (1991); Lin *et al.*, *Science* 254:1022-1024 (1991)); Family III, the metabotropic glutamate receptor family

(Nakanishi, *Science* 258 597:603 (1992)); Family IV, the cAMP receptor family, important in the chemotaxis and development of *D. discoideum* (Klein *et al.*, *Science* 241:1467-1472 (1988)); and Family V, the fungal mating pheromone receptors such as STE2 (Kurjan, *Annu. Rev. Biochem.* 61:1097-1129 (1992)).

5 There are also a small number of other proteins, which present seven putative hydrophobic segments and appear to be unrelated to GPCRs; they have not been shown to couple to G-proteins. *Drosophila* expresses a photoreceptor-specific protein, bride of sevenless (boss), a seven-transmembrane-segment protein, which has been extensively studied and does not show evidence of being a GPCR (Hart *et al.*, *Proc. Natl. Acad. Sci. USA* 90:5047-5051 (1993)). The gene *frizzled* (*fz*)
10 in *Drosophila* is also thought to be a protein with seven transmembrane segments. Like boss, *fz* has not been shown to couple to G-proteins (Vinson *et al.*, *Nature* 338:263-264 (1989)).

 G proteins represent a family of heterotrimeric proteins composed of α , β and γ subunits, that bind guanine nucleotides. These proteins are usually linked to cell surface receptors, e.g., receptors containing seven transmembrane segments. Following ligand binding to the GPCR, a
15 conformational change is transmitted to the G protein, which causes the α -subunit to exchange a bound GDP molecule for a GTP molecule and to dissociate from the $\beta\gamma$ -subunits. The GTP-bound form of the α -subunit typically functions as an effector-modulating moiety, leading to the production of second messengers, such as cAMP (e.g., by activation of adenylyl cyclase), diacylglycerol or inositol phosphates. Greater than 20 different types of α -subunits are known in
20 humans. These subunits associate with a smaller pool of β and γ subunits. Examples of mammalian G proteins include Gi, Go, Gq, Gs and Gt. G proteins are described extensively in Lodish *et al.*, *Molecular Cell Biology*, (Scientific American Books Inc., New York, N.Y., 1995), the contents of which are incorporated herein by reference. GPCRs, G proteins and G protein-linked effector and second messenger systems have been reviewed in *The G-Protein Linked Receptor Fact*
25 *Book*, Watson *et al.*, eds., Academic Press (1994).

Dopamine receptors

 The understanding of the dopaminergic system relevance in brain function and disease developed several decades ago from three diverse observations following drug treatments. These
30 were the observations that dopamine replacement therapy improved Parkinson's disease symptoms, depletion of dopamine and other catecholamines by reserpine caused depression and antipsychotic drugs blocked dopamine receptors. The finding that the dopamine receptor binding affinities of typical antipsychotic drugs correlate with their clinical potency led to the dopamine overactivity

hypothesis of schizophrenia (Snyder, S.H., *Am J Psychiatry* 133, 197-202 (1976); Seeman, P. and Lee, T., *Science* 188, 1217-9 (1975)). Today, dopamine receptors are crucial targets in the pharmacological therapy of schizophrenia, Parkinson's disease, Tourette's syndrome, tardive dyskinesia and Huntington's disease. The dopaminergic system includes the nigrostriatal, mesocorticolimbic and tuberoinfundibular pathways. The nigrostriatal pathway is part of the striatal motor system and its degeneration leads to Parkinson's disease; the mesocorticolimbic pathway plays a key role in reinforcement and in emotional expression and is the desired site of action of antipsychotic drugs; the tuberoinfundibular pathways regulates prolactin secretion from the pituitary.

Dopamine receptors are members of the G protein coupled receptor superfamily, a large group proteins that share a seven helical membrane-spanning structure and transduce signals through coupling to heterotrimeric guanine nucleotide-binding regulatory proteins (G proteins). Dopamine receptors are classified into subfamilies: D1-like (D1 and D5) and D2-like (D2, D3 and D4) based on their different ligand binding profiles, signal transduction properties, sequence homologies and genomic organizations (Civelli, O., Bunzow, J.R. and Grandy, D.K., *Annu Rev Pharmacol Toxicol* 33, 281-307 (1993)). The D1-like receptors, D1 and D5, stimulate cAMP synthesis through coupling with Gs-like proteins and their genes do not contain introns within their protein coding regions. On the other hand, the D2-like receptors, D2, D3 and D4, inhibit cAMP synthesis through their interaction with Gi-like proteins and share a similar genomic organization which includes introns within their protein coding regions.

Serotonin receptors

Serotonin (5-Hydroxytryptamine; 5-HT) was first isolated from blood serum, where it was shown to promote vasoconstriction (Rapport, M.M., Green, A.A. and Page, I.H., *J Biol Chem* 176, 1243-1251 (1948). Interest on a possible relationship between 5-HT and psychiatric disease was spurred by the observations that hallucinogens such as LSD and psilocybin inhibit the actions of 5-HT on smooth muscle preparations (Gaddum, J.H. and Hameed, K.A., *Br J Pharmacol* 9, 240-248 (1954)). This observation lead to the hypothesis that brain 5-HT activity might be altered in psychiatric disorders (Wooley, D.W. and Shaw, E., *Proc Natl Acad Sci U S A* 40, 228-231 (1954); Gaddum, J.H. and Picarelli, Z.P., *Br J Pharmacol* 12, 323-328 (1957)). This hypothesis was strengthened by the introduction of tricyclic antidepressants and monoamine oxidase inhibitors for the treatment of major depression and the observation that those drugs affected noradrenaline and 5-HT metabolism. Today, drugs acting on the serotonergic system have been proved to be effective

in the pharmacotherapy of psychiatric diseases such as depression, schizophrenia, obsessive-compulsive disorder, panic disorder, generalized anxiety disorder and social phobia as well as migraine, vomiting induced by cancer chemotherapy and gastric motility disorders.

Serotonin receptors represent a very large and diverse family of neurotransmitter receptors.

5 To date thirteen 5-HT receptor proteins coupled to G proteins plus one ligand-gated ion channel receptor (5-HT₃) have been described in mammals. This receptor diversity is thought to reflect serotonin's ancient origin as a neurotransmitter and a hormone as well as the many different roles of 5-HT in mammals. The 5-HT receptors have been classified into seven subfamilies or groups according to their different ligand-binding affinity profiles, molecular structure and intracellular
10 transduction mechanisms (Hoyer, D. et al., *Pharmacol. Rev.* 46, 157-203 (1994)).

GPCRs are a major target for drug/insecticide action and development. Accordingly, it is valuable to the field of pharmaceutical development to identify and characterize previously unknown GPCRs. The present invention advances the state of the art by providing a previously unidentified drosophila GPCRs.

15

Insecticides

About 10,000 species of the more than 1 million species of insects are crop-eating, and of these, approximately 700 species worldwide cause most of the insect damage to man's crops, in the field and in storage.

20 A detailed study of novel proteins from *Drosophila* and invertebrate orthologs thereof, will serve as targets for identifying new members of the known classes of insecticides as well as aiding in the identification of new classes of compounds.

SUMMARY OF THE INVENTION

25 The present invention is based in part on the identification of nucleic acid sequences that encode amino acid sequences of 66 GPCR proteins that are produced by *Drosophila melanogaster* and invertebrate, human and other mammalian orthologs thereof. These unique protein sequences, and nucleic acid sequences that encode these proteins, can be used as models for the development of human therapeutic targets, aid in the identification of therapeutic proteins
30 and serve as targets for the development of human therapeutic and insecticidal agents.

The proteins of the present inventions are GPCRs that participate in signaling pathways. As used herein, a "signaling pathway" refers to the modulation (e.g., stimulation or inhibition) of a cellular function/activity upon the binding of a ligand to the GPCR protein. Examples of such

functions include mobilization of intracellular molecules that participate in a signal transduction pathway, e.g., phosphatidylinositol 4,5-bisphosphate (PIP₂), inositol 1,4,5-triphosphate (IP₃) and adenylate cyclase; polarization of the plasma membrane; production or secretion of molecules; alteration in the structure of a cellular component; cell proliferation, e.g., synthesis of DNA; cell migration; cell differentiation; and cell survival

The response mediated by the receptor protein depends on the type of cell. For example, in some cells, binding of a ligand to the receptor protein may stimulate an activity such as release of compounds, gating of a channel, cellular adhesion, migration, differentiation, etc., through phosphatidylinositol or cyclic AMP metabolism and turnover while in other cells, the binding of the ligand will produce a different result. Regardless of the cellular activity/response modulated by the receptor protein, it is universal that the protein is a GPCR and interacts with G proteins to produce one or more secondary signals, in a variety of intracellular signal transduction pathways, e.g., through phosphatidylinositol or cyclic AMP metabolism and turnover, in a cell.

As used herein, "phosphatidylinositol turnover and metabolism" refers to the molecules involved in the turnover and metabolism of phosphatidylinositol 4,5-bisphosphate (PIP₂) as well as to the activities of these molecules. PIP₂ is a phospholipid found in the cytosolic leaflet of the plasma membrane. Binding of ligand to the receptor activates, in some cells, the plasma-membrane enzyme phospholipase C that in turn can hydrolyze PIP₂ to produce 1,2-diacylglycerol (DAG) and inositol 1,4,5-triphosphate (IP₃). Once formed IP₃ can diffuse to the endoplasmic reticulum surface where it can bind an IP₃ receptor, e.g., a calcium channel protein containing an IP₃ binding site. IP₃ binding can induce opening of the channel, allowing calcium ions to be released into the cytoplasm. IP₃ can also be phosphorylated by a specific kinase to form inositol 1,3,4,5-tetraphosphate (IP₄), a molecule that can cause calcium entry into the cytoplasm from the extracellular medium. IP₃ and IP₄ can subsequently be hydrolyzed very rapidly to the inactive products inositol 1,4-bisphosphate (IP₂) and inositol 1,3,4-triphosphate, respectively. These inactive products can be recycled by the cell to synthesize PIP₂. The other second messenger produced by the hydrolysis of PIP₂, namely 1,2-diacylglycerol (DAG), remains in the cell membrane where it can serve to activate the enzyme protein kinase C. Protein kinase C is usually found soluble in the cytoplasm of the cell, but upon an increase in the intracellular calcium concentration, this enzyme can move to the plasma membrane where it can be activated by DAG. The activation of protein kinase C in different cells results in various cellular responses such as the phosphorylation of glycogen synthase, or the phosphorylation of various transcription factors, e.g., NF-kB. The language "phosphatidylinositol activity", as used herein, refers to an activity of PIP₂ or one of its metabolites.

Another signaling pathway in which the receptor may participate is the cAMP turnover pathway. As used herein, "cyclic AMP turnover and metabolism" refers to the molecules involved in the turnover and metabolism of cyclic AMP (cAMP) as well as to the activities of these molecules. Cyclic AMP is a second messenger produced in response to ligand-induced stimulation of certain G protein coupled receptors. In the cAMP signaling pathway, binding of a ligand to a GPCR can lead to the activation of the enzyme adenylyl cyclase, which catalyzes the synthesis of cAMP. The newly synthesized cAMP can in turn activate a cAMP-dependent protein kinase. This activated kinase can phosphorylate a voltage-gated potassium channel protein, or an associated protein, and lead to the inability of the potassium channel to open during an action potential. The inability of the potassium channel to open results in a decrease in the outward flow of potassium, which normally repolarizes the membrane of a neuron, leading to prolonged membrane depolarization.

By targeting an agent to modulate a GPCR, the signaling activity and biological process mediated by the receptor can be agonized or antagonized. Such agonism and antagonism serves as a basis for modulating a biological activity in a therapeutic context (mammalian therapy) or toxic context (anti-invertebrate/insecticidal agent).

DESCRIPTION OF THE FIGURE SHEETS

FIGURE SHEETS 1-89 provides genomic nucleic acid sequences from *Drosophila melanogaster*, predicted transcript, amino acid coding regions, information relating to the subfamily of GPCR proteins to which the protein sequence belongs, e.g., BLAST hit homology, presence of hidden Markov model, etc. Also provided are 5' promoter sequences, transcription initiation sites and other structural features.

DETAILED DESCRIPTION OF THE INVENTION

General Description

The present invention is based on the sequencing of the *Drosophila melanogaster* genome. During the sequencing and assembly of the *Drosophila melanogaster* genome, analysis of the sequence information revealed previously unidentified nucleic acid molecules that encode proteins that share structural and/or sequence homology to protein/peptide/domains identified and characterized within the art as being a GPCR protein or part of a GPCR protein. Based on this analysis, the present invention provides amino acid sequences of 66 GPCR proteins that are produced by *Drosophila melanogaster*, nucleic acid sequences that encode these GPCR proteins,

and the art known protein/peptide/domain that have structural or sequence homology to the *Drosophila melanogaster* protein.

In addition to being previously unknown, the proteins and nucleic acid molecules that are provided in the present invention are selected based on their ability to be used for the development of commercially important products and services. Specifically, the present proteins are selected based on homology and/or structural relatedness to known GPCR proteins of commercial importance and/or the presence of genetic signals that define the protein as a GPCR protein. Some of the more specific features of the proteins of the present invention, and the uses thereof, are described in detail below, and are known within the art for each category of GPCR.

Specific Embodiments

Peptide Molecules

In the Figure sheets, the present invention provides nucleic acid sequences that encode 66 protein molecules that have been identified as being within the *Drosophila melanogaster* genome (SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, encoded by nucleic acid molecules provided in SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)). These protein sequences will be referred herein as the GPCR proteins of the present invention, GPCR proteins, or the peptides or the proteins of the present invention. Table 1 provides a summary of the GPCRs of the present invention and the subfamily assignment of each one.

The present invention provides isolated peptide and protein molecules that consist of, consist essentially of or are comprised of the amino acid sequences of the GPCR proteins disclosed in the Figure sheets (SEQ ID NO: 3, 6, 9, 189, 192, 195, 198, encoded by nucleic acid molecules provided in SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)), as well as all obvious variants of these proteins that are within the art to make and use. Some of these variants are described in detail below.

As used herein, a protein is said to be "isolated" or "purified" when it is substantially free of cellular material or free of chemical precursors or other chemicals. The proteins of the present invention can be purified to homogeneity or other degrees of purity. The level of purification will be based on the intended use. The critical feature is that the preparation allows for the desired function of the protein, even if in the presence of considerable amounts of other components.

In some uses, "substantially free of cellular material" includes preparations of the protein having less than about 30% (by dry weight) other proteins (i.e., contaminating protein), less than about 20% other proteins, less than about 10% other proteins, or less than about 5% other proteins. When the protein is recombinantly produced, it can also be substantially free of culture medium, i.e., culture medium represents less than about 20% of the volume of the protein preparation.

The language "substantially free of chemical precursors or other chemicals" includes preparations of the protein in which it is separated from chemical precursors or other chemicals that are involved in its synthesis. In one embodiment, the language "substantially free of chemical precursors or other chemicals" includes preparations of the GPCR protein having less than about 30% (by dry weight) chemical precursors or other chemicals, less than about 20% chemical precursors or other chemicals, less than about 10% chemical precursors or other chemicals, or less than about 5% chemical precursors or other chemicals.

The isolated GPCR protein can be purified from cells that naturally express it, purified from cells that have been altered to express it (recombinant), or synthesized using known protein synthesis methods. For example, a nucleic acid molecule encoding the GPCR protein is cloned into an expression vector, the expression vector introduced into a host cell and the protein expressed in the host cell. The protein can then be isolated from the cells by an appropriate purification scheme using standard protein purification techniques. Many of these techniques are described in detail below.

Accordingly, the present invention provides proteins that consist of one of the amino acid sequences encoded by the nucleic acid sequences shown in the Figure sheets (SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)). The amino acid sequences of such proteins are provided in the Figure Sheets along with an explanation of the protein type/family. A protein consists of an amino acid sequence when the amino acid sequence is the final amino acid sequence of the protein.

The present invention further provides proteins that consist essentially of one of the amino acid sequences encoded by the nucleic acid sequences shown in the Figure sheets (SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)). A protein consists essentially of an amino acid sequence when such an amino acid sequence is present with only a few additional amino acid residues in the final protein.

The present invention further provides proteins that are comprised of one of the amino acid sequences encoded by the nucleic acid sequences shown in the Figure sheets (SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript

sequences)). A protein is comprised of an amino acid sequence when the amino acid sequence is at least part of the final amino acid sequence of the protein. In such a fashion, the protein can be only the protein or have additional amino acid molecules, such as amino acid residues (contiguous encoded sequence) that are naturally associated with it or heterologous amino acid residues/peptide sequences. Such a protein can have a few additional amino acid residues or can comprise several hundred or more additional amino acids. The preferred classes of proteins that are comprised of the GPCR proteins of the present invention are the naturally occurring mature proteins.

The GPCR proteins of the present invention can be attached to heterologous sequences to form chimeric or fusion proteins. Such chimeric and fusion proteins comprise a GPCR protein operatively linked to a heterologous protein having an amino acid sequence not substantially homologous to the GPCR protein. "Operatively linked" indicates that the GPCR protein and the heterologous protein are fused in-frame. The heterologous protein can be fused to the N-terminus or C-terminus of the GPCR protein.

In some uses, the fusion protein does not affect the activity of the GPCR protein *per se*. For example, the fusion protein can include, but is not limited to, enzymatic fusion proteins, for example beta-galactosidase fusions, yeast two-hybrid GAL fusions, poly-His fusions, MYC-tagged, HI-tagged and Ig fusions. Such fusion proteins, particularly poly-His fusions, can facilitate the purification of recombinant GPCR protein. In certain host cells (e.g., mammalian host cells), expression and/or secretion of a protein can be increased by using a heterologous signal sequence.

A chimeric or fusion protein can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different protein sequences are ligated together in-frame in accordance with conventional techniques. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers which give rise to complementary overhangs between two consecutive gene fragments which can subsequently be annealed and re-amplified to generate a chimeric gene sequence (see Ausubel *et al.*, *Current Protocols in Molecular Biology*, 1992). Moreover, many expression vectors are commercially available that already encode a fusion moiety (e.g., a GST protein). A GPCR protein-encoding nucleic acid can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the GPCR protein.

As mentioned above, the present invention also provides and enables obvious variants of the amino acid sequence of the proteins of the present invention, such as naturally occurring mature forms of the protein, allelic/sequence variants of the proteins, non-naturally occurring

recombinantly derived variants of the proteins, and orthologs and paralogs of the proteins. Such variants can readily be generated using art know techniques in the fields of recombinant nucleic acid technology and protein biochemistry. It is understood, however, that variants exclude any amino acid sequences disclosed prior to the invention.

5 Such variants can readily be identified/made using molecular techniques and the sequence information disclosed herein. Further, such variants can readily be distinguished from other proteins based on sequence and/or structural homology to the GPCR proteins of the present invention. The degree of homology/identity present will be based primarily on whether the protein is a functional variant or non-functional variant, the amount of divergence present in the paralog
10 family and the evolutionary distance between the orthologs.

To determine the percent identity of two amino acid sequences or two nucleic acid sequences, the sequences are aligned for optimal comparison purposes (e.g., gaps can be introduced in one or both of a first and a second amino acid or nucleic acid sequence for optimal alignment and non-homologous sequences can be disregarded for comparison purposes). In a
15 preferred embodiment, the length of a reference sequence aligned for comparison purposes is at least 30%, 40%, 50%, 60%, 70%, 80%, or 90% or more of the length of the reference sequence. The amino acid residues or nucleotides at corresponding amino acid positions or nucleotide positions are then compared. When a position in the first sequence is occupied by the same amino acid residue or nucleotide as the corresponding position in the second sequence, then the
20 molecules are identical at that position (as used herein amino acid or nucleic acid "identity" is equivalent to amino acid or nucleic acid "homology"). The percent identity between the two sequences is a function of the number of identical positions shared by the sequences, taking into account the number of gaps, and the length of each gap, which need to be introduced for optimal alignment of the two sequences.

25 The comparison of sequences and determination of percent identity and similarity between two sequences can be accomplished using a mathematical algorithm. (*Computational Molecular Biology*, Lesk, A.M., ed., Oxford University Press, New York, 1988; *Biocomputing: Informatics and Genome Projects*, Smith, D.W., ed., Academic Press, New York, 1993; *Computer Analysis of Sequence Data, Part 1*, Griffin, A.M., and Griffin, H.G., eds., Humana Press, New
30 Jersey, 1994; *Sequence Analysis in Molecular Biology*, von Heinje, G., Academic Press, 1987; and *Sequence Analysis Primer*, Gribskov, M. and Devereux, J., eds., M Stockton Press, New York, 1991). In a preferred embodiment, the percent identity between two amino acid sequences is determined using the Needleman and Wunsch (*J. Mol. Biol.* (48):444-453 (1970)) algorithm

which has been incorporated into the GAP program in the GCG software package (available at <http://www.gcg.com>), using either a Blossum 62 matrix or a PAM250 matrix, and a gap weight of 16, 14, 12, 10, 8, 6, or 4 and a length weight of 1, 2, 3, 4, 5, or 6. In yet another preferred embodiment, the percent identity between two nucleotide sequences is determined using the
5 GAP program in the GCG software package (Devereux, J., *et al.*, *Nucleic Acids Res.* 12(1):387 (1984)) (available at <http://www.gcg.com>), using a NWSgapdna.CMP matrix and a gap weight of 40, 50, 60, 70, or 80 and a length weight of 1, 2, 3, 4, 5, or 6. In another embodiment, the percent identity between two amino acid or nucleotide sequences is determined using the algorithm of E. Meyers and W. Miller (CABIOS, 4:11-17 (1989)) which has been incorporated
10 into the ALIGN program (version 2.0), using a PAM120 weight residue table, a gap length penalty of 12 and a gap penalty of 4.

The nucleic acid and protein sequences of the present invention can further be used as a "query sequence" to perform a search against sequence databases to, for example, identify other family members or related sequences. Such searches can be performed using the NBLAST and
15 XBLAST programs (version 2.0) of Altschul, et al. (*J. Mol. Biol.* 215:403-10 (1990)). BLAST nucleotide searches can be performed with the NBLAST program, score = 100, wordlength = 12 to obtain nucleotide sequences homologous to the nucleic acid molecules of the invention. BLAST protein searches can be performed with the XBLAST program, score = 50, wordlength = 3 to obtain amino acid sequences homologous to the proteins of the invention. To obtain gapped
20 alignments for comparison purposes, Gapped BLAST can be utilized as described in Altschul et al. (*Nucleic Acids Res.* 25(17):3389-3402 (1997)). When utilizing BLAST and gapped BLAST programs, the default parameters of the respective programs (e.g., XBLAST and NBLAST) can be used. See <http://www.ncbi.nlm.nih.gov>.

Full-length pre-processed forms, as well as mature processed forms, of proteins that
25 comprise one of the proteins of the present invention can readily be identified as having complete sequence identity to one of the GPCR proteins of the present invention as well as being encoded by the same genetic locus as the GPCR protein provided herein.

Allelic variants of a GPCR protein can readily be identified as having a high degree (significant) of sequence homology/identity to at least a portion of the GPCR protein as well as
30 being encoded by the same genetic locus as the GPCR protein provided herein. As used herein, two proteins (or a region of the proteins) have significant homology when the amino acid sequences are typically at least about 70-75%, 80-85%, and more typically at least about 90-95% or more homologous. A significantly homologous amino acid sequence, according to the present

invention, will be encoded by a nucleic acid sequence that will hybridize to a GPCR protein encoding nucleic acid molecule under stringent conditions as more fully described below.

Paralogs of a GPCR protein can readily be identified as having some degree of significant sequence homology/identity to at least a portion of the GPCR protein, as being encoded by a gene
5 from *Drosophila*, and as having similar activity or function. Two proteins will typically be considered paralogs when the amino acid sequences are typically at least about 70-75%, 80-85%, and more typically at least about 90-95% or more homologous through a given region or domain. Such paralogs will be encoded by a nucleic acid sequence that will hybridize to a GPCR protein encoding nucleic acid molecule under stringent conditions as more fully described below.

10 Orthologs of a GPCR protein can readily be identified as having some degree of significant sequence homology/identity to at least a portion of the GPCR protein as well as being encoded by a gene from another organism. Preferred orthologs will be isolated from mammals, preferably human, for the development of human therapeutic targets and agents, or other invertebrates, particularly insects of economical/agriculture importance, e.g. members of the Lepidopteran and
15 Coleopteran orders, for the development of insecticides and insecticidal targets. Such orthologs will be encoded by a nucleic acid sequence that will hybridize to a GPCR protein encoding nucleic acid molecule under moderate to stringent conditions, as more fully described below, depending on the degree of relatedness of the two organisms yielding the proteins.

Non-naturally occurring variants of the GPCR proteins of the present invention can readily
20 be generated using recombinant techniques. Such variants include, but are not limited to deletions, additions and substitutions in the amino acid sequence of the GPCR protein. For example, one class of substitutions are conserved amino acid substitution. Such substitutions are those that substitute a given amino acid in a GPCR protein by another amino acid of like characteristics. Typically seen as conservative substitutions are the replacements, one for another, among the aliphatic amino acids
25 Ala, Val, Leu, and Ile; interchange of the hydroxyl residues Ser and Thr, exchange of the acidic residues Asp and Glu, substitution between the amide residues Asn and Gln, exchange of the basic residues Lys and Arg and replacements among the aromatic residues Phe, Tyr. Guidance concerning which amino acid changes are likely to be phenotypically silent are found in Bowie *et al.*, *Science* 247:1306-1310 (1990).

30 Variant GPCR proteins can be fully functional or can lack function in one or more activities, e.g. ability to bind ligand, ability to bind G-protein, ability to mediate signaling, etc. Fully functional variants typically contain only conservative variation or variation in non-critical residues or in non-critical regions. Functional variants can also contain substitution of similar amino acids

that result in no change or an insignificant change in function. Alternatively, such substitutions may positively or negatively affect function to some degree.

Non-functional variants typically contain one or more non-conservative amino acid substitutions, deletions, insertions, inversions, or truncation or a substitution, insertion, inversion, or deletion in a critical residue or critical region.

Amino acids that are essential for function can be identified by methods known in the art, such as site-directed mutagenesis or alanine-scanning mutagenesis (Cunningham *et al.*, *Science* 244:1081-1085 (1989)). The latter procedure introduces single alanine mutations at every residue in the molecule. The resulting mutant molecules are then tested for biological activity such as receptor binding or in assays such as an *in vitro* proliferative activity. Sites that are critical for ligand-receptor binding can also be determined by structural analysis such as crystallization, nuclear magnetic resonance or photoaffinity labeling (Smith *et al.*, *J. Mol. Biol.* 224:899-904 (1992); de Vos *et al.* *Science* 255:306-312 (1992)).

Polypeptides often contain amino acids other than the 20 amino acids commonly referred to as the 20 naturally-occurring amino acids. Further, many amino acids, including the terminal amino acids, may be modified by natural processes, such as processing and other post-translational modifications, or by chemical modification techniques well known in the art. Common modifications that occur naturally in polypeptides are described in basic texts, detailed monographs, and the research literature, and they are well known to those of skill in the art.

Accordingly, the polypeptides also encompass derivatives or analogs in which a substituted amino acid residue is not one encoded by the genetic code, in which a substituent group is included, in which the mature polypeptide is fused with another compound, such as a compound to increase the half-life of the polypeptide (for example, polyethylene glycol), or in which the additional amino acids are fused to the mature polypeptide, such as a leader or secretory sequence or a sequence for purification of the mature polypeptide or a pro-protein sequence.

The present invention further provides fragments of the GPCR proteins, in addition to proteins and peptides that comprise and consist of such fragments. The fragments to which the invention pertains, however, are not to be construed as encompassing fragments that may be disclosed publicly prior to the present invention.

As used herein, a fragment comprises at least 8 or more contiguous amino acid residues from a GPCR protein. Such fragments can be chosen based on the ability to retain one or more of the biological activities of the GPCR protein or could be chosen for the ability to perform a function, e.g. act as an immunogen. Particularly important fragments are biologically active

fragments, peptides which are, for example, about 8 or more amino acids in length. Such fragments will typically comprise a domain or motif of the GPCR protein, e.g., active site, a G-protein binding site, a transmembrane domain or a ligand binding domain. Further, possible fragments include, but are not limited to, domain or motif containing fragments, soluble peptide fragments, and fragments
5 containing immunogenic structures. Predicted domains and functional sites are readily identifiable by computer programs well-known and readily available to those of skill in the art (e.g., PROSITE analysis).

Polypeptides often contain amino acids other than the 20 amino acids commonly referred to as the 20 naturally-occurring amino acids. Further, many amino acids, including the terminal amino
10 acids, may be modified by natural processes, such as processing and other post-translational modifications, or by chemical modification techniques well known in the art. Common modifications that occur naturally in GPCR proteins are described in basic texts, detailed monographs, and the research literature, and they are well known to those of skill in the art.

Accordingly, the GPCR proteins of the present invention also encompass derivatives or
15 analogs in which a substituted amino acid residue is not one encoded by the genetic code, in which a substituent group is included, in which the mature GPCR protein is fused with another compound, such as a compound to increase the half-life of the GPCR protein (for example, polyethylene glycol), or in which the additional amino acids are fused to the mature GPCR protein, such as a leader or secretory sequence or a sequence for purification of the mature GPCR protein or a pro-
20 protein sequence.

Known modifications include, but are not limited to, acetylation, acylation, ADP-
ribosylation, amidation, covalent attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid
25 derivative, covalent attachment of phosphatidylinositol, cross-linking, cyclization, disulfide bond formation, demethylation, formation of covalent crosslinks, formation of cystine, formation of pyroglutamate, formylation, gamma carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination.

Such modifications are well-known to those of skill in the art and have been described in
30 great detail in the scientific literature. Several particularly common modifications, glycosylation, lipid attachment, sulfation, gamma-carboxylation of glutamic acid residues, hydroxylation and ADP-ribosylation, for instance, are described in most basic texts, such as *Proteins - Structure and*

Molecular Properties, 2nd Ed., T.E. Creighton, W. H. Freeman and Company, New York (1993). Many detailed reviews are available on this subject, such as by Wold, F., *Posttranslational Covalent Modification of Proteins*, B.C. Johnson, Ed., Academic Press, New York 1-12 (1983); Seifter *et al.* (*Meth. Enzymol.* 182: 626-646 (1990)) and Rattan *et al.* (*Ann. N.Y. Acad. Sci.* 663:48-62 (1992)).

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Protein/Peptide Uses

The proteins of the present invention can be used in assays to identify modulators as potential insecticides, to determine the biological activity of the protein, including in a panel of multiple proteins for high-throughput screening; to raise antibodies or to elicit another immune response; as a reagent (including the labeled reagent) in assays designed to quantitatively
10 determine levels of the protein (or its binding partner or receptor) in biological fluids; and as markers for tissues in which the corresponding protein is preferentially expressed. Where the protein binds or potentially binds to another protein (such as, for example, in a receptor-ligand interaction), the protein can be used to identify the binding partner so as to develop a system to
15 identify inhibitors of the binding interaction. Any or all of these research utilities are capable of being developed into reagent grade or kit format for commercialization as research products.

Methods for performing the uses listed above are well known to those skilled in the art. References disclosing such methods include "Molecular Cloning: A Laboratory Manual", 2d ed., Cold Spring Harbor Laboratory Press, Sambrook, J., E. F. Fritsch and T. Maniatis eds., 1989,
20 and "Methods in Enzymology: Guide to Molecular Cloning Techniques", Academic Press, Berger, S. L. and A. R. Kimmel eds., 1987 and can readily be determined using the information provided in Table 1.

The potential uses of the proteins of the present invention are based primarily on the source of the protein as well as the class/action of the protein. For example, GPCRs isolated
25 from *Drosophila* and other invertebrates serve as a target for identifying anti-invertebrate compounds, e.g. insecticides. GPCRs isolated from *Drosophila* and their human/mammalian orthologs serve as targets for identifying agents for use in mammalian therapeutic applications, e.g. a human drug. Approximately 70% of all pharmaceutical agents modulate the activity of a GPCR. A combination of the invertebrate and mammalian ortholog can be used in selective
30 screening methods to find agents specific for invertebrates. Table 1 provides specific context of use for each of the presently disclosed GPCRs. Particularly useful are GPCRs involved in neurotransmission for use as insecticide targets.

The receptor polypeptides (including variants and fragments which may have been disclosed prior to the present invention) are useful for biological assays related to GPCRs. Such assays involve any of the known GPCR functions or activities or properties useful for diagnosis and treatment of GPCR-related conditions.

5 The receptor polypeptides are also useful in drug/insecticide screening assays, in cell-based or cell-free systems. Cell-based systems can be native, i.e., cells that normally express the receptor protein, as a biopsy or expanded in cell culture. In one embodiment, however, cell-based assays involve recombinant host cells expressing the receptor protein.

10 The polypeptides can be used to identify compounds that modulate receptor activity. Both the GPCRs of the present invention and appropriate variants and fragments can be used in high-throughput screens to assay candidate compounds for the ability to bind to the receptor. These compounds can be further screened against a functional receptor to determine the effect of the compound on the receptor activity. Further, these compounds can be tested in animal or invertebrate systems to determine activity/effectiveness. Compounds can be identified that activate
15 (agonist) or inactivate (antagonist) the receptor to a desired degree.

Further, the receptor polypeptides can be used to screen a compound for the ability to stimulate or inhibit interaction between the receptor protein and a target molecule that normally interacts with the receptor protein. The target can be ligand or a component of the signal pathway that the receptor protein normally interacts (for example, a G-protein or other interactor involved in
20 cAMP or phosphatidylinositol turnover and/or adenylate cyclase, or phospholipase C activation). Such assays typically include the steps of combining the receptor protein with a candidate compound under conditions that allow the receptor protein, or fragment, to interact with the target molecule, and to detect the formation of a complex between the protein and the target or to detect the biochemical consequence of the interaction with the receptor protein and the target, such as any
25 of the associated effects of signal transduction such as G-protein phosphorylation, cAMP or phosphatidylinositol turnover, and adenylate cyclase or phospholipase C activation.

Candidate compounds include, for example, 1) peptides such as soluble peptides, including Ig-tailed fusion peptides and members of random peptide libraries (see, e.g., Lam *et al.*, *Nature* 354:82-84 (1991); Houghten *et al.*, *Nature* 354:84-86 (1991)) and combinatorial chemistry-derived
30 molecular libraries made of D- and/or L- configuration amino acids; 2) phosphopeptides (e.g., members of random and partially degenerate, directed phosphopeptide libraries, see, e.g., Songyang *et al.*, *Cell* 72:767-778 (1993)); 3) antibodies (e.g., polyclonal, monoclonal, humanized, anti-idiotypic, chimeric, and single chain antibodies as well as Fab, F(ab')₂, Fab expression library

fragments, and epitope-binding fragments of antibodies); and 4) small organic and inorganic molecules (e.g., molecules obtained from combinatorial and natural product libraries).

One candidate compound is a soluble fragment of the receptor that competes for ligand binding. Other candidate compounds include mutant receptors or appropriate fragments containing mutations that affect receptor function and thus compete for ligand. Accordingly, a fragment that competes for ligand, for example with a higher affinity, or a fragment that binds ligand but does not allow release, is encompassed by the invention.

The invention further includes other end point assays to identify compounds that modulate (stimulate or inhibit) receptor activity. The assays typically involve an assay of events in the signal transduction pathway that indicate receptor activity. Thus, the expression of genes that are up- or down-regulated in response to the receptor protein dependent signal cascade can be assayed. In one embodiment, the regulatory region of such genes can be operably linked to a marker that is easily detectable, such as luciferase. Alternatively, phosphorylation of the receptor protein, or a receptor protein target, could also be measured.

Any of the biological or biochemical functions mediated by the receptor can be used as an endpoint assay. These include all of the biochemical or biochemical/biological events described herein, in the references cited herein, incorporated by reference for these endpoint assay targets, and other functions known to those of ordinary skill in the art.

Binding and/or activating compounds can also be screened by using chimeric receptor proteins in which the amino terminal extracellular domain, or parts thereof, the entire transmembrane domain or subregions, such as any of the seven transmembrane segments or any of the intracellular or extracellular loops and the carboxy terminal intracellular domain, or parts thereof, can be replaced by heterologous domains or subregions. For example, a G-protein-binding region can be used that interacts with a different G-protein than that which is recognized by the native receptor. Accordingly, a different set of signal transduction components is available as an end-point assay for activation. Alternatively, the entire transmembrane portion or subregions (such as transmembrane segments or intracellular or extracellular loops) can be replaced with the entire transmembrane portion or subregions specific to a host cell that is different from the host cell from which the amino terminal extracellular domain and/or the G-protein-binding region are derived. This allows for assays to be performed in other than the specific host cell from which the receptor is derived. Alternatively, the amino terminal extracellular domain (and/or other ligand-binding regions) could be replaced by a domain (and/or other binding region) binding a different ligand, thus, providing an assay for test compounds that interact with the heterologous amino terminal

extracellular domain (or region) but still cause signal transduction. Finally, activation can be detected by a reporter gene containing an easily detectable coding region operably linked to a transcriptional regulatory sequence that is part of the native signal transduction pathway.

The receptor polypeptides are also useful in competition binding assays in methods designed to discover compounds that interact with the receptor. Thus, a compound is exposed to a receptor polypeptide under conditions that allow the compound to bind or to otherwise interact with the polypeptide. Soluble receptor polypeptide is also added to the mixture. If the test compound interacts with the soluble receptor polypeptide, it decreases the amount of complex formed or activity from the receptor target. This type of assay is particularly useful in cases in which compounds are sought that interact with specific regions of the receptor. Thus, the soluble polypeptide that competes with the target receptor region is designed to contain peptide sequences corresponding to the region of interest.

To perform cell free drug/insecticide screening assays, it is sometimes desirable to immobilize either the receptor protein, or fragment, or its target molecule to facilitate separation of complexes from uncomplexed forms of one or both of the proteins, as well as to accommodate automation of the assay.

Techniques for immobilizing proteins on matrices can be used in the drug/insecticide screening assays. In one embodiment, a fusion protein can be provided which adds a domain that allows the protein to be bound to a matrix. For example, glutathione-S-transferase/15625 fusion proteins can be adsorbed onto glutathione sepharose beads (Sigma Chemical, St. Louis, MO) or glutathione derivatized microtitre plates, which are then combined with the cell lysates (e.g., ³⁵S-labeled) and the candidate compound, and the mixture incubated under conditions conducive to complex formation (e.g., at physiological conditions for salt and pH). Following incubation, the beads are washed to remove any unbound label, and the matrix immobilized and radiolabel determined directly, or in the supernatant after the complexes are dissociated. Alternatively, the complexes can be dissociated from the matrix, separated by SDS-PAGE, and the level of receptor-binding protein found in the bead fraction quantitated from the gel using standard electrophoretic techniques. For example, either the polypeptide or its target molecule can be immobilized utilizing conjugation of biotin and streptavidin using techniques well known in the art. Alternatively, antibodies reactive with the protein but which do not interfere with binding of the protein to its target molecule can be derivatized to the wells of the plate, and the protein trapped in the wells by antibody conjugation. Preparations of a receptor-binding protein and a candidate compound are incubated in the receptor protein-presenting wells and the amount of complex trapped in the well

can be quantitated. Methods for detecting such complexes, in addition to those described above for the GST-immobilized complexes, include immunodetection of complexes using antibodies reactive with the receptor protein target molecule, or which are reactive with receptor protein and compete with the target molecule, as well as enzyme-linked assays which rely on detecting an enzymatic activity associated with the target molecule.

Agents that modulate one of the GPCRs of the present invention can be identified using one or more of the above assays, alone or in combination. It is generally preferable to use a cell-based or cell free system first and then confirm activity in an animal/insect model system. Such model systems are well known in the art and can readily be employed in this context.

Modulators of receptor protein activity identified according to these drug/insecticide screening assays can be used to treat a subject with a disorder mediated by the receptor pathway, by treating cells that express the GPCR. These methods of treatment include the steps of administering the modulators of protein activity in a pharmaceutical composition as described herein, to a subject in need of such treatment.

In yet another aspect of the invention, the GPCR proteins can be used as "bait proteins" in a two-hybrid assay or three-hybrid assay (see, e.g., U.S. Patent No. 5,283,317; Zervos et al. (1993) *Cell* 72:223-232; Madura et al. (1993) *J. Biol. Chem.* 268:12046-12054; Bartel et al. (1993) *Biotechniques* 14:920-924; Iwabuchi et al. (1993) *Oncogene* 8:1693-1696; and Brent WO94/10300), to identify other proteins, which bind to or interact with the GPCR and are involved in GPCR activity. Such GPCR-binding proteins are also likely to be involved in the propagation of signals by the GPCR proteins or GPCR targets as, for example, downstream elements of a GPCR-mediated signaling pathway, e.g., a signaling pathway. Alternatively, such GPCR-binding proteins are likely to be GPCR inhibitors.

The two-hybrid system is based on the modular nature of most transcription factors, which consist of separable DNA-binding and activation domains. Briefly, the assay utilizes two different DNA constructs. In one construct, the gene that codes for a GPCR protein is fused to a gene encoding the DNA binding domain of a known transcription factor (e.g., GAL-4). In the other construct, a DNA sequence, from a library of DNA sequences, that encodes an unidentified protein ("prey" or "sample") is fused to a gene that codes for the activation domain of the known transcription factor. If the "bait" and the "prey" proteins are able to interact, *in vivo*, forming a GPCR-dependent complex, the DNA-binding and activation domains of the transcription factor are brought into close proximity. This proximity allows transcription of a reporter gene (e.g., LacZ) which is operably linked to a transcriptional regulatory site responsive to the transcription

factor. Expression of the reporter gene can be detected and cell colonies containing the functional transcription factor can be isolated and used to obtain the cloned gene which encodes the protein which interacts with the GPCR protein.

This invention further pertains to novel agents identified by the above-described screening assays. Accordingly, it is within the scope of this invention to further use an agent identified as described herein in an appropriate animal model. For example, an agent identified as described herein (e.g., a GPCR modulating agent, an antisense GPCR nucleic acid molecule, a GPCR-specific antibody, or a GPCR-binding partner) can be used in an animal or insect model to determine the efficacy, toxicity, or side effects of treatment with such an agent. Alternatively, an agent identified as described herein can be used in an animal or insect model to determine the mechanism of action of such an agent. Furthermore, this invention pertains to uses of novel agents identified by the above-described screening assays for insecticidal activity as described herein.

15 Antibodies

The invention also provides antibodies that selectively bind to one of the proteins of the present invention, a protein comprising such a peptide, as well as variants and fragments thereof. As used herein, an antibody selectively binds a target peptide when it binds the target peptide and does not significantly bind to unrelated proteins. An antibody is still considered to selectively bind a peptide even if it also binds to other proteins that are not substantially homologous with the target peptide so long as such proteins share homology with a fragment or domain of the peptide target of the antibody. In this case, it would be understood that antibody binding to the peptide is still selective despite some degree of cross-reactivity.

As used herein, an antibody is defined in terms consistent with that recognized within the art: they are multi-subunit proteins produced by a mammalian organism in response to an antigen challenge. The antibodies of the present invention include polyclonal antibodies and monoclonal antibodies, as well as fragments of such antibodies, including, but not limited to, Fab or F(ab')₂, and Fv fragments.

Many methods are known for generating and/or identifying antibodies to a given target peptide. Several such methods are described by Harlow, Antibodies, Cold Spring Harbor Press, (1989).

In general, to generate antibodies, an isolated peptide is used as an immunogen and is administered to a mammalian organism, such as a rat, rabbit or mouse. Either the full-length protein, an antigenic peptide fragment or a fusion protein can be used.

Antibodies are preferably prepared from regions or discrete fragments of the GPCR proteins. Antibodies can be prepared from any region of the protein as described herein. However, preferred regions will include those involved in function/activity and/or receptor/binding partner interaction.

An antigenic fragment will typically comprise at least 10 contiguous amino acid residues. The antigenic peptide can comprise, however, at least 12, 14, 20 or more amino acid residues. Such fragments can be selected on a physical property, such as fragments correspond to regions that are located on the surface of the protein, e.g., hydrophilic regions or can be selected based on sequence uniqueness.

Detection on an antibody of the present invention can be facilitated by coupling (i.e., physically linking) the antibody to a detectable substance. Examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, and radioactive materials. Examples of suitable enzymes include horseradish peroxidase, alkaline phosphatase, β -galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin, and examples of suitable radioactive material include ^{125}I , ^{131}I , ^{35}S or ^3H .

Antibody Uses

The antibodies can be used to isolate one of the proteins of the present invention by standard techniques, such as affinity chromatography or immunoprecipitation. The antibodies can facilitate the purification of the natural protein from cells and recombinantly produced protein expressed in host cells. In addition, such antibodies are useful to detect the presence of one of the proteins of the present invention in cells or tissues to determine the pattern of expression of the protein among various tissues in an organism and over the course of normal development. Further, such antibodies can be used to detect protein *in situ*, *in vitro*, or in a cell lysate or supernatant in order to evaluate the abundance and pattern of expression. Also, such antibodies can be used to assess abnormal tissue

distribution or abnormal expression during development. Antibody detection of circulating fragments of the full length protein can be used to identify turnover.

The antibodies can also be used to assess normal and aberrant subcellular localization of cells in the various tissues in an organism. The diagnostic uses can be applied, not only in genetic testing, but also in monitoring a treatment modality. Accordingly, where treatment is ultimately aimed at correcting expression level or the presence of aberrant sequence and aberrant tissue distribution or developmental expression, antibodies directed against the or relevant fragments can be used to monitor therapeutic efficacy.

The antibodies are also useful for inhibiting protein function, for example, blocking the binding of the GPCR protein to a binding partner such as a ligand. An antibody can be used, for example, to block binding, thus modulating (agonizing or antagonizing) the peptides activity. Antibodies can be prepared against specific fragments containing sites required for function or against intact protein that is associated with a cell or cell membrane.

The invention also encompasses kits for using antibodies to detect the presence of a protein in a biological sample. The kit can comprise antibodies such as a labeled or labelable antibody and a compound or agent for detecting protein in a biological sample; means for determining the amount of protein in the sample; means for comparing the amount of protein in the sample with a standard; and instructions for use.

Nucleic Acid Molecules

The present invention further provides isolated nucleic acid molecules that encode a GPCR protein or protein of the present invention (SEQ ID NO: 3, 6, 9, . . . , 192, 195, 198, encoded by nucleic acid molecules provided in SEQ ID NOS: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOS: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)). Such nucleic acid molecules will consist of, consist essentially of, or comprise a nucleotide sequence that encodes one of the GPCR proteins of the present invention, an allelic variant thereof, or an ortholog or paralog thereof.

As used herein, an "isolated" nucleic acid molecule is one that is separated from other nucleic acid present in the natural source of the nucleic acid. Preferably, an "isolated" nucleic acid is free of sequences which naturally flank the nucleic acid (i.e., sequences located at the 5' and 3' ends of the nucleic acid) in the genomic DNA of the organism from which the nucleic acid is derived. However, there can be some flanking nucleotide sequences, for example up to about 5KB, 4KB, 3KB, 2KB, or 1KB or less, particularly contiguous protein encoding sequences and protein

encoding sequences within the same gene but separated by introns in the genomic sequence. The important point is that the nucleic acid is isolated from remote and unimportant flanking sequences such that it can be subjected to the specific manipulations described herein such as recombinant expression, preparation of probes and primers, and other uses specific to the nucleic acid sequences.

5 Moreover, an "isolated" nucleic acid molecule, such as a cDNA molecule, can be substantially free of other cellular material, or culture medium when produced by recombinant techniques, or chemical precursors or other chemicals when chemically synthesized. However, the nucleic acid molecule can be fused to other coding or regulatory sequences and still be considered isolated.

10 For example, recombinant DNA molecules contained in a vector are considered isolated. Further examples of isolated DNA molecules include recombinant DNA molecules maintained in heterologous host cells or purified (partially or substantially) DNA molecules in solution. Isolated RNA molecules include *in vivo* or *in vitro* RNA transcripts of the isolated DNA molecules of the present invention. Isolated nucleic acid molecules according to the present invention further include
15 such molecules produced synthetically.

 Accordingly, the present invention provides nucleic acid molecules that consist of one of the nucleotide sequences shown in the Figure sheets (SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)). A nucleic acid molecule consists of a nucleotide sequence when the nucleotide sequence is the
20 complete nucleotide sequence of the nucleic acid molecule.

 The present invention further provides nucleic acid molecules that consist essentially of one of the nucleotide sequences shown in the Figure sheets (SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)). A nucleic acid molecule consists essentially of a nucleotide sequence when such a nucleotide
25 sequence is present with only a few additional nucleic acid residues in the final nucleic acid molecule.

 The present invention further provides nucleic acid molecules that are comprised of one of the nucleotide sequences shown in the Figure sheets (SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)). A
30 nucleic acid molecule is comprised of a nucleotide sequence when the nucleotide sequence is at least part of the final nucleotide sequence of the nucleic acid molecule. In such a fashion, the nucleic acid molecule can be only the nucleotide sequence or have additional nucleic acid residues, such as nucleic acid residues that are naturally associated with it or heterologous nucleotide

sequences. Such a nucleic acid molecule can have a few additional nucleotides or can comprises several hundred or more additional nucleotides. The preferred classes of nucleic acid molecules that are comprised of the nucleotide sequences of the present are the naturally occurring full-length cDNA molecules and genes and genomic clones since some of the nucleic acid molecules provided
5 in the Figure sheets are fragments of the complete gene that exists in nature. A brief description of how various types of these nucleic acid molecules can be readily made/isolated is provided below.

In the Figures, both coding and non-coding sequences are provided for each protein encoding nucleic acid sequence. Because of the source of the present invention, *Drosophila* genomic sequences, the nucleic acid molecules in the figures will contain genomic intronic
10 sequences, 5' and 3' non-coding sequences, gene regulatory regions and non-coding intergenic sequences. In general such sequence features are either noted or can readily be identified using computational tools known in the art. As discussed below, some of the non-coding regions, particularly gene regulatory elements such as promoters, are useful for a variety of purposes, e.g. control of heterologous gene expression, target for identifying gene activity modulating
15 compounds.

Full-length genes may be cloned from known sequence using any one of a number of methods known in the art. For example, a method which employs XL-PCR (Perkin-Elmer, Foster City, Calif.) to amplify long pieces of DNA may be used. Other methods for obtaining full-length sequences are well known in the art.

The isolated nucleic acid molecules can encode the mature protein plus additional amino or carboxyl-terminal amino acids, or amino acids interior to the mature protein (when the mature form has more than one protein chain, for instance). Such sequences may play a role in processing of a protein from precursor to a mature form, facilitate protein trafficking, prolong or shorten protein half-life or facilitate manipulation of a protein for assay or production, among other things. As
20 generally is the case *in situ*, the additional amino acids may be processed away from the mature protein by cellular enzymes.

As mentioned above, the isolated nucleic acid molecules include, but are not limited to, the sequence encoding the GPCR protein alone, the sequence encoding the mature protein and additional coding sequences, such as a leader or secretory sequence (e.g., a pre-pro or pro-protein
30 sequence), the sequence encoding the mature protein, with or without the additional coding sequences, plus additional non-coding sequences, for example introns and non-coding 5' and 3' sequences such as transcribed but non-translated sequences that play a role in transcription, mRNA processing (including splicing and polyadenylation signals), ribosome binding and stability of

mRNA. In addition, the nucleic acid molecule may be fused to a marker sequence encoding, for example, a protein that facilitates purification.

Isolated nucleic acid molecules can be in the form of RNA, such as mRNA, or in the form DNA, including cDNA and genomic DNA obtained by cloning or produced by chemical synthetic techniques or by a combination thereof. The nucleic acid, especially DNA, can be double-stranded or single-stranded. Single-stranded nucleic acid can be the coding strand (sense strand) or the non-coding strand (anti-sense strand).

The invention further provides nucleic acid molecules that encode fragments of the proteins of the present invention as well as nucleic acid molecules that encode obvious variants of the GPCR proteins of the present invention that are described above. Such nucleic acid molecules may be naturally occurring, such as allelic variants (same locus), paralogs (different locus), and orthologs (different organism), or may be constructed by recombinant DNA methods or by chemical synthesis. Such non-naturally occurring variants may be made by mutagenesis techniques, including those applied to nucleic acid molecules, cells, or organisms. Accordingly, as discussed above, the variants can contain nucleotide substitutions, deletions, inversions and insertions. Variation can occur in either or both the coding and non-coding regions. The variations can produce both conservative and non-conservative amino acid substitutions.

The present invention further provides non-coding fragments of the nucleic acid molecules provided in the Figures. Preferred non-coding fragments include, but are not limited to, promoter sequences, enhancer sequences, gene modulating sequences and gene termination sequences. Such fragments are useful in controlling heterologous gene expression and in developing screens to identify gene modulating agents. Particularly useful are fragments from about 100 to about 1,000 nucleotides taken 5' from the start ATG in the genomic sequences provided in the Figures (SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196,).

A fragment comprises a contiguous nucleotide sequence greater than 12 or more nucleotides. Further, a fragment could be at least 30, 40, 50, 100, 250 or 500 nucleotides in length. The length of the fragment will be based on its intended use. For example, the fragment can encode epitope bearing regions of the protein, or can be useful as DNA probes and primers. Such fragments can be isolated using the known nucleotide sequence to synthesize an oligonucleotide probe. A labeled probe can then be used to screen a cDNA library, genomic DNA library, or mRNA to isolate nucleic acid corresponding to the coding region. Further, primers can be used in PCR reactions to clone specific regions of gene.

A probe/primer typically comprises substantially a purified oligonucleotide or oligonucleotide pair. The oligonucleotide typically comprises a region of nucleotide sequence that hybridizes under stringent conditions to at least about 12, 20, 25, 40, 50 or more consecutive nucleotides.

5 Orthologs, homologs, and allelic variants can be identified using methods well known in the art. As described in the protein Section, these variants comprise a nucleotide sequence encoding a protein that is typically 60-65%, 70-75%, 80-85%, and more typically at least about 90-95% or more homologous to the nucleotide sequence shown in the Figure sheets or a fragment of this sequence. Such nucleic acid molecules can readily be identified as being able to hybridize under
10 moderate to stringent conditions, to the nucleotide sequence shown in the Figure sheets or a fragment of the sequence.

As used herein, the term "hybridizes under stringent conditions" is intended to describe conditions for hybridization and washing under which nucleotide sequences encoding a protein at least 50-55% homologous to each other typically remain hybridized to each other. The conditions
15 can be such that sequences at least about 65%, at least about 70%, or at least about 75% or more homologous to each other typically remain hybridized to each other. Such stringent conditions are known to those skilled in the art and can be found in *Current Protocols in Molecular Biology*, John Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6. One example of stringent hybridization conditions are hybridization in 6X sodium chloride/sodium citrate (SSC) at about 45C, followed by one or more
20 washes in 0.2 X SSC, 0.1% SDS at 50-65C.

Nucleic Acid Molecule Uses

The nucleic acid molecules of the present invention are useful for probes, primers, chemical intermediates, and in biological assays. The nucleic acid molecules are useful as a hybridization
25 probe for cDNA and genomic DNA to isolate full-length cDNA and genomic clones encoding the protein described in the Figures and to isolate cDNA and genomic clones that correspond to variants (alleles, orthologs, etc.) producing the same or related proteins shown in the Figures.

The probe can correspond to any sequence along the entire length of the nucleic acid molecules provided in the Figures. Accordingly, it could be derived from 5' noncoding regions, the
30 coding region, and 3' noncoding regions. However, as discussed, fragments are not to be construed as encompassing fragments disclosed prior to the present invention.

The nucleic acid molecules are also useful as primers for PCR to amplify any given region of a nucleic acid molecule and are useful to synthesize antisense molecules of desired length and sequence.

The nucleic acid molecules are also useful for constructing recombinant vectors. Such
5 vectors include expression vectors that express a portion of, or all of, the protein sequences. Vectors also include insertion vectors, used to integrate into another nucleic acid molecule sequence, such as into the cellular genome, to alter *in situ* expression of a gene and/or gene product. For example, an endogenous coding sequence can be replaced via homologous recombination with all or part of the coding region containing one or more specifically introduced mutations.

10 The nucleic acid molecules are also useful for expressing antigenic portions of the proteins.

The nucleic acid molecules are also useful in making vectors containing the gene regulatory regions of the nucleic acid molecules of the present invention.

The nucleic acid molecules are also useful for designing ribozymes corresponding to all, or a part, of the mRNA produced from the nucleic acid molecules described herein.

15 The nucleic acid molecules are also useful for constructing host cells expressing a part, or all, of the nucleic acid molecules and proteins.

The nucleic acid molecules are also useful for constructing transgenic animals expressing all, or a part, of the nucleic acid molecules and proteins.

20 The nucleic acid molecules are also useful for making vectors that express part, or all, of the proteins.

The nucleic acid molecules are also useful as hybridization probes for determining the presence, level, form and distribution of nucleic acid expression. Accordingly, the probes can be used to detect the presence of, or to determine levels of, a specific nucleic acid molecule in cells, tissues, and in organisms. The nucleic acid whose level is determined can be DNA or RNA.

25 Accordingly, probes corresponding to the proteins described herein can be used to assess expression and/or gene copy number in a given cell, tissue, or organism. These uses are relevant for efficacy of insecticides involving an increase or decrease in GPCR protein expression relative to normal results.

In vitro techniques for detection of mRNA include Northern hybridizations and *in situ* hybridizations. *In vitro* techniques for detecting DNA includes Southern hybridizations and *in situ*
30 hybridization.

Probes can be used as a part of a diagnostic test kit for identifying cells or tissues that express a GPCR protein, such as by measuring a level of a receptor-encoding nucleic acid in a

sample of cells from a subject e.g., mRNA or genomic DNA, or determining if a receptor gene has been mutated.

Nucleic acid expression assays are useful for drug/insecticide screening to identify compounds that modulate GPCR nucleic acid expression.

5 The invention thus provides a method for identifying a compound that can be used to treat a block the activity of the GPCR by modulating the expression of the GPCR gene. The method typically includes assaying the ability of the compound to modulate the expression of the GPCR nucleic acid and thus identifying a compound that can be used to kill insects by altering GPCR nucleic acid expression. The assays can be performed in cell-based and cell-free systems. Cell-
10 based assays include cells naturally expressing the GPCR nucleic acid or recombinant cells genetically engineered to express specific nucleic acid sequences.

The assay for GPCR nucleic acid expression can involve direct assay of nucleic acid levels, such as mRNA levels, or on collateral compounds involved in the signal pathway. Further, the expression of genes that are up- or down-regulated in response to the GPCR protein signal pathway
15 can also be assayed. In this embodiment the regulatory regions of these genes can be operably linked to a reporter gene such as luciferase.

Thus, modulators of GPCR gene expression can be identified in a method wherein a cell is contacted with a candidate compound and the expression of mRNA determined. The level of expression of GPCR mRNA in the presence of the candidate compound is compared to the level of
20 expression of GPCR mRNA in the absence of the candidate compound. The candidate compound can then be identified as a modulator of nucleic acid expression based on this comparison and be used, for example to treat a disorder characterized by aberrant nucleic acid expression. When expression of mRNA is statistically significantly greater in the presence of the candidate compound than in its absence, the candidate compound is identified as a stimulator of nucleic acid expression.
25 When nucleic acid expression is statistically significantly less in the presence of the candidate compound than in its absence, the candidate compound is identified as an inhibitor of nucleic acid expression.

The nucleic acid molecules are also useful in diagnostic assays for qualitative changes in GPCR nucleic acid, and particularly in qualitative changes that lead to insecticidal
30 activity/tolerance. The nucleic acid molecules can be used to detect mutations in GPCR genes and gene expression products such as mRNA.

The nucleic acid molecules are thus useful as antisense constructs to control GPCR gene expression in cells, tissues, and organisms. A DNA antisense nucleic acid molecule is designed to

be complementary to a region of the gene involved in transcription, preventing transcription and hence production of GPCR protein. An antisense RNA or DNA nucleic acid molecule would hybridize to the mRNA and thus block translation of mRNA into GPCR protein.

5 The invention also encompasses kits for detecting the presence of a GPCR nucleic acid in a biological sample. For example, the kit can comprise reagents such as a labeled or labelable nucleic acid or agent capable of detecting GPCR nucleic acid in a biological sample; means for determining the amount of GPCR nucleic acid in the sample; and means for comparing the amount of GPCR nucleic acid in the sample with a standard. The compound or agent can be packaged in a suitable container. The kit can further comprise instructions for using the kit to detect GPCR protein mRNA
10 or DNA.

Vectors/host cells

The invention also provides vectors containing the nucleic acid molecules described herein. The term "vector" refers to a vehicle, preferably a nucleic acid molecule, that can transport the
15 nucleic acid molecules. When the vector is a nucleic acid molecule, the nucleic acid molecules are covalently linked to the vector nucleic acid. With this aspect of the invention, the vector includes a plasmid, single or double stranded phage, a single or double stranded RNA or DNA viral vector, or artificial chromosome, such as a BAC, PAC, YAC, OR MAC.

A vector can be maintained in the host cell as an extrachromosomal element where it
20 replicates and produces additional copies of the nucleic acid molecules. Alternatively, the vector may integrate into the host cell genome and produce additional copies of the nucleic acid molecules when the host cell replicates.

The invention provides vectors for the maintenance (cloning vectors) or vectors for expression (expression vectors) of the nucleic acid molecules. The vectors can function in
25 procaryotic or eukaryotic cells or in both (shuttle vectors).

Expression vectors contain cis-acting regulatory regions that are operably linked in the vector to the nucleic acid molecules such that transcription of the nucleic acid molecules is allowed in a host cell. The nucleic acid molecules can be introduced into the host cell with a separate nucleic acid molecule capable of affecting transcription. Thus, the second nucleic acid molecule
30 may provide a trans-acting factor interacting with the cis-regulatory control region to allow transcription of the nucleic acid molecules from the vector. Alternatively, a trans-acting factor may be supplied by the host cell. Finally, a trans-acting factor can be produced from the vector itself. It

is understood, however, that in some embodiments, transcription and/or translation of the nucleic acid molecules can occur in a cell-free system.

The regulatory sequence to which the nucleic acid molecules described herein can be operably linked include promoters for directing mRNA transcription. These include, but are not limited to, the left promoter from bacteriophage λ , the lac, TRP, and TAC promoters from *E. coli*, the early and late promoters from SV40, the CMV immediate early promoter, the adenovirus early and late promoters, and retrovirus long-terminal repeats.

In addition to control regions that promote transcription, expression vectors may also include regions that modulate transcription, such as repressor binding sites and enhancers.

Examples include the SV40 enhancer, the cytomegalovirus immediate early enhancer, polyoma enhancer, adenovirus enhancers, and retrovirus LTR enhancers.

In addition to containing sites for transcription initiation and control, expression vectors can also contain sequences necessary for transcription termination and, in the transcribed region a ribosome binding site for translation. Other regulatory control elements for expression include initiation and termination codons as well as polyadenylation signals. The person of ordinary skill in the art would be aware of the numerous regulatory sequences that are useful in expression vectors. Such regulatory sequences are described, for example, in Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual*. 2nd. ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, (1989).

A variety of expression vectors can be used to express a nucleic acid molecule. Such vectors include chromosomal, episomal, and virus-derived vectors, for example vectors derived from bacterial plasmids, from bacteriophage, from yeast episomes, from yeast chromosomal elements, including yeast artificial chromosomes, from viruses such as baculoviruses, papovaviruses such as SV40, Vaccinia viruses, adenoviruses, poxviruses, pseudorabies viruses, and retroviruses. Vectors may also be derived from combinations of these sources such as those derived from plasmid and bacteriophage genetic elements, e.g. cosmids and phagemids. Appropriate cloning and expression vectors for prokaryotic and eukaryotic hosts are described in Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual*. 2nd. ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, (1989).

The regulatory sequence may provide constitutive expression in one or more host cells (i.e. tissue specific) or may provide for inducible expression in one or more cell types such as by temperature, nutrient additive, or exogenous factor such as a hormone or other ligand. A variety of

vectors providing for constitutive and inducible expression in prokaryotic and eukaryotic hosts are well known to those of ordinary skill in the art.

The nucleic acid molecules can be inserted into the vector nucleic acid by well-known methodology. Generally, the DNA sequence that will ultimately be expressed is joined to an expression vector by cleaving the DNA sequence and the expression vector with one or more
5 restriction enzymes and then ligating the fragments together. Procedures for restriction enzyme digestion and ligation are well known to those of ordinary skill in the art.

The vector containing the appropriate nucleic acid molecule can be introduced into an appropriate host cell for propagation or expression using well-known techniques. Bacterial cells
10 include, but are not limited to, *E. coli*, *Streptomyces*, and *Salmonella typhimurium*. Eukaryotic cells include, but are not limited to, yeast, insect cells such as *Drosophila*, animal cells such as COS and CHO cells, and plant cells.

As described herein, it may be desirable to express the protein as a fusion protein. Accordingly, the invention provides fusion vectors that allow for the production of the proteins.
15 Fusion vectors can increase the expression of a recombinant protein, increase the solubility of the recombinant protein, and aid in the purification of the protein by acting for example as a ligand for affinity purification. A proteolytic cleavage site may be introduced at the junction of the fusion moiety so that the desired peptide can ultimately be separated from the fusion moiety. Proteolytic enzymes include, but are not limited to, factor Xa, thrombin, and enterokinase. Typical fusion
20 expression vectors include pGEX (Smith *et al.*, *Gene* 67:31-40 (1988)), pMAL (New England Biolabs, Beverly, MA) and pRIT5 (Pharmacia, Piscataway, NJ) which fuse glutathione S-transferase (GST), maltose E binding protein, or protein A, respectively, to the target recombinant protein. Examples of suitable inducible non-fusion *E. coli* expression vectors include pTrc (Amann
25 *et al.*, *Gene* 69:301-315 (1988)) and pET 11d (Studier *et al.*, *Gene Expression Technology: Methods in Enzymology* 185:60-89 (1990)).

Recombinant protein expression can be maximized in a host bacteria by providing a genetic background wherein the host cell has an impaired capacity to proteolytically cleave the recombinant protein. (Gottesman, S., *Gene Expression Technology: Methods in Enzymology* 185, Academic Press, San Diego, California (1990) 119-128). Alternatively, the sequence of the nucleic acid
30 molecule of interest can be altered to provide preferential codon usage for a specific host cell, for example *E. coli*. (Wada *et al.*, *Nucleic Acids Res.* 20:2111-2118 (1992)).

The nucleic acid molecules can also be expressed by expression vectors that are operative in yeast. Examples of vectors for expression in yeast e.g., *S. cerevisiae* include pYepSec1 (Baldari, *et*

al., *EMBO J.* 6:229-234 (1987)), pMFa (Kurjan *et al.*, *Cell* 30:933-943(1982)), pJRY88 (Schultz *et al.*, *Gene* 54:113-123 (1987)), and pYES2 (Invitrogen Corporation, San Diego, CA).

The nucleic acid molecules can also be expressed in insect cells using, for example, baculovirus expression vectors. Baculovirus vectors available for expression of proteins in cultured
5 insect cells (e.g., Sf 9 cells) include the pAc series (Smith *et al.*, *Mol. Cell Biol.* 3:2156-2165 (1983)) and the pVL series (Lucklow *et al.*, *Virology* 170:31-39 (1989)).

In certain embodiments of the invention, the nucleic acid molecules described herein are expressed in mammalian cells using mammalian expression vectors. Examples of mammalian expression vectors include pCDM8 (Seed, B. *Nature* 329:840(1987)) and pMT2PC (Kaufman *et al.*,
10 *EMBO J.* 6:187-195 (1987)).

The expression vectors listed herein are provided by way of example only of the well-known vectors available to those of ordinary skill in the art that would be useful to express the nucleic acid molecules. The person of ordinary skill in the art would be aware of other vectors suitable for maintenance propagation or expression of the nucleic acid molecules described herein.
15 These are found for example in Sambrook, J., Fritsh, E. F., and Maniatis, T. *Molecular Cloning: A Laboratory Manual*. 2nd, ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, 1989.

The invention also encompasses vectors in which the nucleic acid sequences described herein are cloned into the vector in reverse orientation, but operably linked to a regulatory sequence
20 that permits transcription of antisense RNA. Thus, an antisense transcript can be produced to all, or to a portion, of the nucleic acid molecule sequences described herein, including both coding and non-coding regions. Expression of this antisense RNA is subject to each of the parameters described above in relation to expression of the sense RNA (regulatory sequences, constitutive or inducible expression, tissue-specific expression).

25 The invention also relates to recombinant host cells containing the vectors described herein. Host cells therefore include prokaryotic cells, lower eukaryotic cells such as yeast, other eukaryotic cells such as insect cells, and higher eukaryotic cells such as mammalian cells.

The recombinant host cells are prepared by introducing the vector constructs described herein into the cells by techniques readily available to the person of ordinary skill in the art. These
30 include, but are not limited to, calcium phosphate transfection, DEAE-dextran-mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection, lipofection, and other techniques such as those found in Sambrook, *et al.* (*Molecular Cloning: A*

Laboratory Manual. 2nd, ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, 1989).

Host cells can contain more than one vector. Thus, different nucleotide sequences can be introduced on different vectors of the same cell. Similarly, the nucleic acid molecules can be introduced either alone or with other nucleic acid molecules that are not related to the nucleic acid molecules such as those providing trans-acting factors for expression vectors. When more than one vector is introduced into a cell, the vectors can be introduced independently, co-introduced or joined to the nucleic acid molecule vector.

In the case of bacteriophage and viral vectors, these can be introduced into cells as packaged or encapsulated virus by standard procedures for infection and transduction. Viral vectors can be replication-competent or replication-defective. In the case in which viral replication is defective, replication will occur in host cells providing functions that complement the defects.

Vectors generally include selectable markers that enable the selection of the subpopulation of cells that contain the recombinant vector constructs. The marker can be contained in the same vector that contains the nucleic acid molecules described herein or may be on a separate vector. Markers include tetracycline or ampicillin-resistance genes for prokaryotic host cells and dihydrofolate reductase or neomycin resistance for eukaryotic host cells. However, any marker that provides selection for a phenotypic trait will be effective.

While the mature proteins can be produced in bacteria, yeast, mammalian cells, and other cells under the control of the appropriate regulatory sequences, cell-free transcription and translation systems can also be used to produce these proteins using RNA derived from the DNA constructs described herein.

Where secretion of the protein is desired, which is difficult to achieve with multi-transmembrane domain containing proteins such as GPCRs, appropriate secretion signals are incorporated into the vector. The signal sequence can be endogenous to the proteins or heterologous to these proteins.

Where the protein is not secreted into the medium, which is typically the case with GPCRs, the protein can be isolated from the host cell by standard disruption procedures, including freeze thaw, sonication, mechanical disruption, use of lysing agents and the like. The protein can then be recovered and purified by well-known purification methods including ammonium sulfate precipitation, acid extraction, anion or cationic exchange chromatography, phosphocellulose chromatography, hydrophobic-interaction chromatography, affinity chromatography,

hydroxylapatite chromatography, lectin chromatography, or high performance liquid chromatography.

It is also understood that depending upon the host cell in recombinant production of the proteins described herein, the proteins can have various glycosylation patterns, depending upon the cell, or maybe non-glycosylated as when produced in bacteria. In addition, the proteins may include an initial modified methionine in some cases as a result of a host-mediated process.

Nucleic Acid Arrays

The present invention further provides arrays or microarrays of nucleic acid molecules that are based on the sequence information provided in the Figure Sheets SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)).

As used herein "Arrays" or "Microarrays" refers to an array of distinct polynucleotides or oligonucleotides synthesized on a substrate, such as paper, nylon or other type of membrane, filter, chip, glass slide, or any other suitable solid support. In one embodiment, the microarray is prepared and used according to the methods described in US Patent 5,837,832, Chee et al., PCT application W095/11995 (Chee et al.), Lockhart, D. J. et al. (1996; Nat. Biotech. 14: 1675-1680) and Schena, M. et al. (1996; Proc. Natl. Acad. Sci. 93: 10614-10619), all of which are incorporated herein in their entirety by reference. In other embodiments, such arrays are produced by the methods described by Brown et. al., US Patent No. 5,807,522.

The microarray is preferably composed of a large number of unique, single-stranded nucleic acid sequences, usually either synthetic antisense oligonucleotides or fragments of cDNAs, fixed to a solid support. The oligonucleotides are preferably about 6-60 nucleotides in length, more preferably 15-30 nucleotides in length, and most preferably about 20-25 nucleotides in length. For a certain type of microarray, it may be preferable to use oligonucleotides that are only 7-20 nucleotides in length. The microarray may contain oligonucleotides that cover the known 5', or 3', sequence, sequential oligonucleotides which cover the full length sequence; or unique oligonucleotides selected from particular areas along the length of the sequence. Polynucleotides used in the microarray may be oligonucleotides that are specific to a gene or genes of interest.

In order to produce oligonucleotides to a known sequence for a microarray, the gene(s) of interest (or an ORF identified from the contigs of the present invention) is typically examined using a computer algorithm which starts at the 5' or at the 3' end of the nucleotide sequence.

Typical algorithms will then identify oligomers of defined length that are unique to the gene, have a GC content within a range suitable for hybridization, and lack predicted secondary structure that may interfere with hybridization. In certain situations it may be appropriate to use pairs of oligonucleotides on a microarray. The "pairs" will be identical, except for one
5 nucleotide that preferably is located in the center of the sequence. The second oligonucleotide in the pair (mismatched by one) serves as a control. The number of oligonucleotide pairs may range from two to one million. The oligomers are synthesized at designated areas on a substrate using a light-directed chemical process. The substrate may be paper, nylon or other type of membrane, filter, chip, glass slide or any other suitable solid support.

10 In another aspect, an oligonucleotide may be synthesized on the surface of the substrate by using a chemical coupling procedure and an ink jet application apparatus, as described in PCT application W095/251116 (Baldeschweiler et al.) which is incorporated herein in its entirety by reference. In another aspect, a "gridded" array analogous to a dot (or slot) blot may be used to arrange and link cDNA fragments or oligonucleotides to the surface of a substrate using a
15 vacuum system, thermal, UV, mechanical or chemical bonding procedures. An array, such as those described above, may be produced by hand or by using available devices (slot blot or dot blot apparatus), materials (any suitable solid support), and machines (including robotic instruments), and may contain 8, 24, 96, 384, 1536, 6144 or more oligonucleotides, or any other number between two and one million which lends itself to the efficient use of commercially
20 available instrumentation.

In order to conduct sample analysis using a microarray, the RNA or DNA from a biological sample is made into hybridization probes. The mRNA is isolated, and cDNA is produced and used as a template to make antisense RNA (aRNA). The aRNA is amplified in the presence of fluorescent nucleotides, and labeled probes are incubated with the microarray so that
25 the probe sequences hybridize to complementary oligonucleotides of the microarray. Incubation conditions are adjusted so that hybridization occurs with precise complementary matches or with various degrees of less complementarity. After removal of nonhybridized probes, a scanner is used to determine the levels and patterns of fluorescence. The scanned images are examined to determine degree of complementarity and the relative abundance of each oligonucleotide
30 sequence on the microarray. The biological samples may be obtained from any bodily fluids (such as blood, urine, saliva, phlegm, gastric juices, etc.), cultured cells, biopsies, or other tissue preparations. A detection system may be used to measure the absence, presence, and amount of hybridization for all of the distinct sequences simultaneously. This data may be used for large

scale correlation studies on the sequences, expression patterns, mutations, variants, or polymorphisms among samples.

Using such arrays, the present invention provides methods to identify the expression of one or more of the proteins/peptides of the present invention. In detail, such methods comprise
5 incubating a test sample with one or more nucleic acid molecules and assaying for binding of the nucleic acid molecule with components within the test sample. Such assays will typically involve arrays comprising many genes, at least one of which is a gene of the present invention.

Conditions for incubating a nucleic acid molecule with a test sample vary. Incubation conditions depend on the format employed in the assay, the detection methods employed, and the
10 type and nature of the nucleic acid molecule used in the assay. One skilled in the art will recognize that any one of the commonly available hybridization, amplification or array assay formats can readily be adapted to employ the novel fragments of the *Drosophila* genome disclosed herein. Examples of such assays can be found in Chard, T, *An Introduction to Radioimmunoassay and Related Techniques*, Elsevier Science Publishers, Amsterdam, The
15 Netherlands (1986); Bullock, G. R. *et al.*, *Techniques in Immunocytochemistry*, Academic Press, Orlando, FL Vol. 1 (1982), Vol. 2 (1983), Vol. 3 (1985); Tijssen, P., *Practice and Theory of Enzyme Immunoassays: Laboratory Techniques in Biochemistry and Molecular Biology*, Elsevier Science Publishers, Amsterdam, The Netherlands (1985).

The test samples of the present invention include cells, protein or membrane extracts of
20 cells. The test sample used in the above-described method will vary based on the assay format, nature of the detection method and the tissues, cells or extracts used as the sample to be assayed. Methods for preparing nucleic acid extracts or of cells are well known in the art and can be readily be adapted in order to obtain a sample that is compatible with the system utilized.

In another embodiment of the present invention, kits are provided which contain the
25 necessary reagents to carry out the assays of the present invention.

Specifically, the invention provides a compartmentalized kit to receive, in close confinement, one or more containers which comprises: (a) a first container comprising one of the nucleic acid molecules that can bind to a fragment of the *Drosophila* genome disclosed herein; and (b) one or more other containers comprising one or more of the following: wash reagents,
30 reagents capable of detecting presence of a bound nucleic acid. Preferred kits will include chips that are capable of detecting the expression of 10 or more, or 50 or more, or all of the GPCR genes expressed in *Drosophila*.

In detail, a compartmentalized kit includes any kit in which reagents are contained in separate containers. Such containers include small glass containers, plastic containers, strips of plastic, glass or paper, or arraying material such as silica. Such containers allows one to efficiently transfer reagents from one compartment to another compartment such that the samples and reagents are not cross-contaminated, and the agents or solutions of each container can be added in a quantitative fashion from one compartment to another. Such containers will include a container which will accept the test sample, a container which contains the nucleic acid probe, containers which contain wash reagents (such as phosphate buffered saline, Tris-buffers, etc.), and containers which contain the reagents used to detect the bound probe. One skilled in the art will readily recognize that the previously unidentified GPCR genes of the present invention can be routinely identified using the sequence information disclosed herein can be readily incorporated into one of the established kit formats which are well known in the art, particularly expression arrays.

Uses of vectors and host cells

The recombinant host cells expressing the proteins described herein have a variety of uses. First, the cells are useful for producing a GPCR protein or protein that can be further purified to produce desired amounts of GPCR protein or fragments. Thus, host cells containing expression vectors are useful for protein production.

Host cells are also useful for conducting cell-based assays involving the GPCR protein or GPCR protein fragments, such as those described above as well as other formats known in the art. Thus, a recombinant host cell expressing a native GPCR protein is useful for assaying compounds that stimulate or inhibit GPCR protein function.

Host cells are also useful for identifying GPCR protein mutants in which these functions are affected. If the mutants naturally occur and give rise to a pathology, host cells containing the mutations are useful to assay compounds that have a desired effect on the mutant GPCR protein (for example, stimulating or inhibiting function) which may not be indicated by their effect on the native GPCR protein.

Genetically engineered host cells can be further used to produce non-human transgenic animals. A transgenic animal is preferably a mammal, for example a rodent, such as a rat or mouse, in which one or more of the cells of the animal include a transgene. A transgene is exogenous DNA which is integrated into the genome of a cell from which a transgenic animal develops and which remains in the genome of the mature animal in one or more cell types or tissues of the transgenic

animal. These animals are useful for studying the function of a GPCR protein and identifying and evaluating modulators of GPCR protein activity. Other examples of transgenic animals include non-human primates, sheep, dogs, cows, goats, chickens, and amphibians.

A transgenic animal can be produced by introducing nucleic acid into the male pronuclei of a fertilized oocyte, e.g., by microinjection, retroviral infection, and allowing the oocyte to develop in a pseudopregnant female foster animal. Any of the GPCR protein nucleotide sequences can be introduced as a transgene into the genome of a non-human animal, such as a mouse.

Any of the regulatory or other sequences useful in expression vectors can form part of the transgenic sequence. This includes intronic sequences and polyadenylation signals, if not already included. A tissue-specific regulatory sequence(s) can be operably linked to the transgene to direct expression of the GPCR protein to particular cells.

Methods for generating transgenic animals via embryo manipulation and microinjection, particularly animals such as mice, have become conventional in the art and are described, for example, in U.S. Patent Nos. 4,736,866 and 4,870,009, both by Leder *et al.*, U.S. Patent No. 4,873,191 by Wagner *et al.* and in Hogan, B., *Manipulating the Mouse Embryo*, (Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1986). Similar methods are used for production of other transgenic animals. A transgenic founder animal can be identified based upon the presence of the transgene in its genome and/or expression of transgenic mRNA in tissues or cells of the animals. A transgenic founder animal can then be used to breed additional animals carrying the transgene. Moreover, transgenic animals carrying a transgene can further be bred to other transgenic animals carrying other transgenes. A transgenic animal also includes animals in which the entire animal or tissues in the animal have been produced using the homologously recombinant host cells described herein.

In another embodiment, transgenic non-human animals can be produced which contain selected systems which allow for regulated expression of the transgene. One example of such a system is the *cre/loxP* recombinase system of bacteriophage P1. For a description of the *cre/loxP* recombinase system, see, e.g., Lakso *et al.* *PNAS* 89:6232-6236 (1992). Another example of a recombinase system is the FLP recombinase system of *S. cerevisiae* (O'Gorman *et al.* *Science* 251:1351-1355 (1991)). If a *cre/loxP* recombinase system is used to regulate expression of the transgene, animals containing transgenes encoding both the *Cre* recombinase and a selected protein is required. Such animals can be provided through the construction of "double" transgenic animals, e.g., by mating two transgenic animals, one containing a transgene encoding a selected protein and the other containing a transgene encoding a recombinase.

Clones of the non-human transgenic animals described herein can also be produced according to the methods described in Wilmut, I. *et al. Nature* 385:810-813 (1997) and PCT International Publication Nos. WO 97/07668 and WO 97/07669. In brief, a cell, e.g., a somatic cell, from the transgenic animal can be isolated and induced to exit the growth cycle and enter G₀ phase.

5 The quiescent cell can then be fused, e.g., through the use of electrical pulses, to an enucleated oocyte from an animal of the same species from which the quiescent cell is isolated. The reconstructed oocyte is then cultured such that it develops to morula or blastocyst and then transferred to pseudopregnant female foster animal. The offspring born of this female foster animal will be a clone of the animal from which the cell, e.g., the somatic cell, is isolated.

10 Transgenic animals containing recombinant cells that express the proteins described herein are useful to conduct the assays described herein in an *in vivo* context. Accordingly, the various physiological factors that are present *in vivo* and that could effect ligand binding, GPCR protein activation, and signal transduction, may not be evident from *in vitro* cell-free or cell-based assays. Accordingly, it is useful to provide non-human transgenic animals to assay *in vivo* GPCR protein
15 function, including ligand interaction, the effect of specific mutant GPCR proteins on GPCR protein function and ligand interaction, and the effect of chimeric GPCR proteins. It is also possible to assess the effect of null mutations, that is mutations that substantially or completely eliminate one or more GPCR protein functions.

All publications and patents mentioned in the above specification are herein incorporated
20 by reference. Various modifications and variations of the described method and system of the invention will be apparent to those skilled in the art without departing from the scope and spirit of the invention. Although the invention has been described in connection with specific preferred embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the above-
25 described modes for carrying out the invention which are obvious to those skilled in the field of molecular biology or related fields are intended to be within the scope of the following claims.

Table 1

Gene	Transcript	Similar to	F	W	V	Gene	Transcript	Similar to	F	W	V
			E	E	E				E	E	E
A. Rhodopsin-like receptor family						Orphan receptors					
Opsin-like						Related to <i>C. elegans</i> orphan receptors					
CG5638	CT17820	Rh3	e-57	e-14	e-44	CG2114	CT2366		>e-10	e-27	e-10
						CG3171	CT10621	EG:22E5.10	e-50	e-16	e-26
						CG5936	CT18637		>e-10	e-18	>e-10
Receptors for biogenic amines and related compounds						CG8985	CT25824	CG13803	e-162	e-31	>e-10
5-HT receptor-like						CG13803	CT33298	CG8985	e-162	e-36	>e-10
CG8007	CT24060	5-HT2	e-21	e-35	e-27	CG13229	CT32473	CG13803	e-58	e-35	e-10
Muscarinic Acetylcholine receptor-like						Other orphan receptors					
CG7918	CT23924	mAcR-60C	e-32	e-49	e-41	CG9569	CT17758		>e-10	>e-10	>e-10
Unclassifiable biogenic amine receptor-like						CG12290	CT19320		>e-10	>e-10	>e-10
CG17004	CT37739	5-HT7	e-18	e-20	e-28	CG6986	CT21642	CG16726	e-10	>e-10	>e-10
CG7431	CT22855	CG16766	e-70	e-45	e-37	CG13579	CT32961		>e-10	e-18	>e-10
CG16766	CT37292	CG7431	e-53	e-31	e-22	CG13995	CT33551		>e-10	e-13	e-11
CG12796	CT38338	CG6919	e-23	e-20	e-30	CG7497	CT23019		>e-10	>e-10	>e-10
CG6919	CT21432	CG6989	e-82	e-46	e-57	Calcitonin receptor-like					
CG6989	CT21650	CG6919	e-82	e-39	e-46	CG4395	CT4121	CG17415	e-46	e-16	e-41
CG7078	CT21843	CG6919	e-73	e-46	e-38	CG17415	CT38445	CG4395	e-42	e-20	e-61
CG18314	CT41076	CG6919	e-21	e-48	e-25	CG13758	CT33238	CG8422	e-40#	e-65	e-66
CG7994	CT24036		>e-10	>e-10	e-10	B. Sex Diuretic hormone receptor-like					
Purine receptors						CG8422	CT24513	CG12370	e-122	e-25	e-65
Adenosine receptor-like						CG12370	CT24959	CG8422	e-122	e-30	e-101
CG9753	CT27563	CG6989	e-19	e-29	e-42	HE6 receptor-like					
Peptide receptors						CG11318	CT31591	CG15556	e-104	e-11	e-20
Allatostatin receptor-like						CG15556	CT35672	CG11318	e-103	>e-11	e-6
CG10001	CT28187	EG:121E7.2	e-62	e-35	e-34	Latrophilin-like					
FSH/TSH/LH receptor-like						CG8639	CT8755		>e-10	e-47	e-45
CG4187	CT13764	CG5042	e-48	e-23	e-24	Methuselah-like					
CG5042	CT16185	CG4187	e-44	e-25	e-23	CG4521	CT14539	CG6965	e-30	e-13	
Gastrin/CCK receptor-like						CG17795	CT16507	mth	e-132	>e-10	>e-10
CG6857	CT21155	CG6881	e-96	e-33	e-19	CG6530	CT20339	CG6536	e-104	>e-10	>e-10
CG6881	CT21314	CG6857	e-96	e-23	e-33	CG6536	CT20351	CG6530	e-115	>e-10	
Gonadotropin releasing hormone receptor-like						CG6965	CT21585	CG17795	e-12	>e-10	e-10
CG10698	CT29989	GRHR	e-43	e-34	e-39	CG7476	CT22963	mth	e-66	>e-10	>e-10
Growth hormone secretagogue receptor-like						CG13406	CT32762	CG17084	e-47	>e-10	>e-10
CG8784	CT25324	CG8795	e-165	e-38	e-35	CG17084	CT33414	CG13406	e-47	>e-10	>e-10
CG8795	CT25350	CG8784	e-165	e-40	e-37	CG17061	CT33415	mth	e-80	>e-10	>e-10
CG9918	CT27924	CG8795	e-83	e-53	e-33	CG16992	CT37715	mth	e-32	>e-10	>e-10
Tachykinin receptor-like						C. Metabotropic glutamate receptor family					
CG1147	CT1960	NepYr	e-31	e-43	e-40	GABA-B receptor-like					
CG10626	CT29768	Takr99D	e-53	e-41	e-44	CG3022	CT9836	CG6706	e-96	e-56	e-85
Somatostatin receptor-like						CG6706	CT20836	CG3022	e-120	e-61	e-169
CG7285	CT22465	CG13702	e-96	e-24	e-51	CG15274	CT35221	CG3022	e-49	e-156	e-129
CG13702	CT33159	CG7285	e-96	e-26	e-46	Metabotropic glutamate receptor-like					
Vasopressin receptor-like						CG8692	CT5032	Glu-RA	e-152	e-115	e-157
CG6111	CT19191	GRHR	e-36	e-26	e-45	Other					
Unclassifiable peptide receptors						CG7155	CT22117		>e-10	>e-10	>e-10
CG13575	CT32957	CG10626	e-08	e-08	e-8#	CG11923	CT35779	CG17215	e-17	>e-10	e-17
CG14003	CT33559	Takr99D	e-14	e-12	e-14#	D. Atypical 7 TM proteins					
CG5911	CT18539	Takr86C	e-18	e-19	e-26#	Frizzled-like					
CG10823	CT18916	CG6857	e-10	e-10	e-12	CG4626	CT9057	Iz	e-36	e-54	e-76

Claims

That which is claimed is:

1. An isolated protein consisting of an amino acid sequence selected from the group consisting of:
 - (a) the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198;
 - (b) the amino acid sequence of an allelic variant of the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said allelic variant is encoded by a nucleic acid molecule that hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 795, 798, 801 (genomic sequences); SEQ ID NOs: 2, 5, 8, . . . , 796, 799, 802 (transcript sequences), under stringent conditions;
 - (c) the amino acid sequence of an ortholog of an amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said ortholog is encoded by a nucleic acid molecule that hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 795, 798, 801 (genomic sequences); SEQ ID NOs: 2, 5, 8, . . . , 796, 799, 802 (transcript sequences), under stringent condition;
 - (d) a fragment of the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein the fragment comprises at least 10 contiguous amino acids.
2. An isolated protein comprising an amino acid sequence selected from the group consisting of:
 - (a) the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198;
 - (b) the amino acid sequence of an allelic variant of the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said allelic variant is encoded by a nucleic acid molecule that hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 795, 798, 801 (genomic sequences); SEQ ID NOs: 2, 5, 8, . . . , 796, 799, 802 (transcript sequences), under stringent conditions;
 - (c) the amino acid sequence of an ortholog of an amino acid sequence shown in the SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said ortholog is encoded by a nucleic acid molecule that hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 795, 798, 801 (genomic sequences); SEQ ID NOs: 2, 5, 8, . . . , 796, 799, 802 (transcript sequences), under stringent condition;

(d) a fragment of the amino acid sequence shown in the SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein the fragment comprises at least 10 contiguous amino acids.

3. An isolated antibody that selectively binds to a protein of claim 1.
4. An isolated nucleic acid molecule consisting of a nucleotide sequence selected from the group consisting of:
 - (a) a nucleotide sequence that encodes the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198;
 - (b) a nucleotide sequence that encodes of an allelic variant of the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said nucleic acid molecule hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)), under stringent conditions;
 - (c) a nucleotide sequence that encodes an ortholog of an amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said nucleic acid molecule hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)), under stringent condition;
 - (d) a nucleotide sequence that encodes a fragment of the amino acid sequence shown in the SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said the fragment comprises at least 10 contiguous amino acids; and
 - (e) A nucleic acid molecule that is the complement of a nucleic acid molecule of (a)-(d).

5. An isolated nucleic acid molecule comprising a nucleotide sequence selected from the group consisting of:
 - (a) a nucleotide sequence that encodes the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198;
 - (b) a nucleotide sequence that encodes of an allelic variant of the amino acid sequence shown in SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said nucleic acid molecule hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196

(genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)), under stringent conditions;

(c) a nucleotide sequence that encodes an ortholog of an amino acid sequence shown in the SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said nucleic acid molecule hybridizes to the nucleic acid molecule shown in SEQ ID NOs: 1, 4, 7, . . . , 190, 193, 196 (genomic sequences) or SEQ ID NOs: 2, 5, 8, . . . , 191, 194, 197 (transcript sequences)), under stringent condition;

(d) a nucleotide sequence that encodes a fragment of the amino acid sequence shown in the SEQ ID NOs: 3, 6, 9, . . . , 192, 195, 198, wherein said the fragment comprises at least 10 contiguous amino acids; and

(f) A nucleic acid molecule that is the complement of a nucleic acid molecule of (a)-(d).

6. A nucleic acid vector comprising the nucleic acid sequences of claim 4.
7. A nucleic acid vector comprising the nucleic acid sequences of claim 5.
8. A host cell containing the vector of claim 6.
9. A host cell containing the vector of claim 7.
10. A method for producing any of the proteins of claim 1 comprising introducing a nucleotide sequence encoding any of the protein sequences in (a)-(d) into a host cell, and culturing the host cell under conditions in which the proteins are expressed from the nucleic acid.
11. A method for producing any of the proteins of claim 2 comprising introducing a nucleotide sequence encoding any of the protein sequences in (a)-(d) into a host cell, and culturing the host cell under conditions in which the proteins are expressed from the nucleic acid.
12. A method for detecting the presence of any of the proteins of claim 1 in a sample, said method comprising contacting said sample with an agent that specifically allows detection of the presence of the protein in the sample and then detecting the presence of the protein.

13. A method for detecting the presence of any of the proteins of claim 2 in a sample, said method comprising contacting said sample with an agent that specifically allows detection of the presence of the protein in the sample and then detecting the presence of the protein.

14. A kit comprising reagents used for the method of claim 12, wherein the reagents comprise an agent that specifically binds to said protein.

15. A method for detecting the presence of a nucleic acid sequence of claim 4 in a sample, the method comprising contacting the sample with an oligonucleotide that hybridizes to the nucleic acid sequences under stringent conditions and determining whether the oligonucleotide binds to the nucleic acid sequence in the sample.

16. A method for detecting the presence of a nucleic acid sequence of claim 5 in a sample, the method comprising contacting the sample with an oligonucleotide that hybridizes to the nucleic acid sequences under stringent conditions and determining whether the oligonucleotide binds to the nucleic acid sequence in the sample.

17. A kit comprising reagents used for the method of claim 15, wherein the reagents comprise a compound that hybridizes under stringent conditions to any of the nucleic acid molecules.

18. A kit comprising reagents used for the method of claim 16, wherein the reagents comprise a compound that hybridizes under stringent conditions to any of the nucleic acid molecules.

19. A method for identifying an agent that binds to any of the proteins of claim 1, said method comprising contacting the protein with an agent and assaying the contacted mixture to determine whether a complex is formed with the agent bound to the protein.

20. A method for identifying an agent that binds to any of the proteins of claim 2, said method comprising contacting the protein with an agent and assaying the contacted mixture to determine whether a complex is formed with the agent bound to the protein.

SEQUENCE LISTING

<110> Cravgn~~ME~~; Anibal

<120> ISOLATED G-PROTEIN COUPLED RECEPTORS,
NUCLEIC ACID MOLECULES ENCODING GPCR PROTEINS, AND USES
THEREOF AS INSECTICIDAL TARGETS

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<210> 24
 <211> 676
 <212> PRT
 <213> Drosophila

<400> 24

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          20            25            30
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41

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Pro His Gly Val Trp Phe Phe Thr Asp Leu Ile Asn Ala Leu Gln Gly		
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Val Phe Ile Phe Ile Val Val Gly Cys Gln Pro Gln Val Trp Thr Ala		
580	585	590
Cys Arg Arg Ile Phe Cys Pro Arg Leu Arg His Asp Ile Thr Asn Thr		
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Thr Asn Gly Val Gln His Ser Ser Ser Gln Gly Leu Pro Ser Met		
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Thr Asn Thr Thr Ala Thr His Met Pro Ser Asn Pro Ala Glu Asp Glu		
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<211> 3187

<212> DNA

<213> Drosophila

<400> 25

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<210> 26

<211> 1080

<212> DNA

<213> Drosophila

<400> 26

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<210> 27

<211> 359

<212> PRT

<213> Drosophila

<400> 27

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          20           25           30

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 115 120 125
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 Tyr Tyr Gly Ser Tyr Ser Gly Thr Cys Phe Pro Leu His Ile His Glu
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 225 230 235 240
 Cys Glu Phe Ala Val Arg Phe Phe Phe Ile Val Leu Thr Asp Phe Leu
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 305 310 315 320
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<211> 3759

<212> DNA

<213> Drosophila

<400> 28

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<210> 29

<211> 1449

<212> DNA

<213> Drosophila

<400> 29


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<210> 30

<211> 482

<212> PRT

<213> Drosophila

<400> 30

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 35          40          45
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Cys Lys Leu Arg Thr Cys Val Arg Phe Cys Cys Pro His Asp His Ile
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Gln Arg His Tyr Lys Lys Glu Leu Met Val Gln Trp Asp Leu Pro Lys
130          135          140
Pro Cys Asp Asp Met Phe Tyr Leu Asp Asn Arg Asp Ile Met Asp Glu
145          150          155          160
Tyr Thr Leu Phe Glu Asn Gly Arg Leu Leu Arg His Tyr Asp Gln Val
165          170          175
Tyr Leu Asp Lys Ser Glu Tyr Cys Leu Gln His Arg Thr Phe Gly Glu
180          185          190
Gly Asn Asn Asn Ser Ile Arg Ile Ile Pro His Asn Cys Leu Ile Leu
195          200          205
Pro Ser Arg Thr Gly Gln Thr Val Val Met Ile Thr Ser Leu Ile Cys
210          215          220

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<211> 4469

<212> DNA

<213> Drosophila

<400> 31

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<211> 1389

<212> DNA

<213> Drosophila

<400> 32

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<211> 462

<212> PRT

<213> Drosophila

<400> 33

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115           120           125
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Gln Gln Leu Ala Gly Gly Gly Ala Ala Gly Ala Gly Asp Gly Gly Asp

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Asn Arg Thr Ala Pro Ser Thr	Thr Ser Tyr Asp	Gly Gly Gly Ser Gly
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Gly Gly Ala Gly Gly Ser Gly	Gly Ser Thr Phe	Met Leu Leu Leu Glu
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<211> 11501

<212> DNA

<213> Drosophila

<400> 34

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<212> DNA

<213> Drosophila

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<212> PRT

<213> Drosophila

<400> 39

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<211> 3821

<212> DNA

<213> Drosophila

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 <211> 1525
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 <213> Drosophila

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<210> 45
 <211> 507
 <212> PRT
 <213> Drosophila

<400> 45
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 50 55 60
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<211> 3688
<212> DNA
<213> Drosophila
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<210> 47

<211> 1191

<212> DNA

<213> Drosophila

<400> 47


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<210> 48

<211> 396

<212> PRT

<213> Drosophila

<400> 48

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165          170          175
Leu Val Ser Ala Thr Leu Phe Ala Ile Pro Ala Leu Ile Ile Ser Ala
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Cys Tyr Ala Ile Ile Val Lys Thr Ile Trp Ala Lys Gly Ser Ile Phe
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210          215          220
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<210> 49

<211> 5051

<212> DNA

<213> Drosophila

<400> 49

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<211> 3051

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<213> Drosophila

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<210> 51

<211> 798

<212> PRT

<213> Drosophila

<400> 51

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74

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Gln Tyr Ser Ser Asn Ser	Cys Lys Tyr Leu Thr	Pro Gln Ser Ser Leu
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Ser Asp Glu Gln Pro Pro	Ala Thr Pro Pro Pro	Thr Val Ala Val Ala
725	730	735
Pro Pro Pro Thr Arg Pro	Gln Arg Pro Lys Leu	Gln Arg Gly Ile Thr
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<211> 3919

<212> DNA

<213> Drosophila

<400> 52

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<210> 53

<211> 1547

<212> DNA

<213> Drosophila

<400> 53

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<211> 192

<212> PRT

<213> Drosophila

<400> 54

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Ala Cys His Leu Arg Pro Cys Ile Arg Phe Cys Cys Pro Gln Tyr Gln
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<211> 3640

<212> DNA

<213> Drosophila

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 <211> 1089
 <212> DNA
 <213> Drosophila

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<210> 57
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 <212> PRT
 <213> Drosophila

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<210> 58

<211> 7503

<212> DNA

<213> Drosophila

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<212> DNA

<213> Drosophila

<400> 62

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<211> 749

<212> PRT

<213> Drosophila

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93

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<213> Drosophila

<400> 64

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<211> 1248

<212> DNA

<213> Drosophila

<400> 65

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 Arg Tyr Leu Lys Phe Ile Ala Asp Gly Leu Ile Asp Glu Gly Leu Gly
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 Ser Ala Val Gly Ser Gly Ser Ser Ile Ala Val Ser Val Glu Asp Val
 65 70 75 80
 Val Ala Gly Gln Ala Gln Asp Ile Gln Ala Ser Glu Gly Ser Thr Asp
 85 90 95
 Asp Ala Asp Gly Ser Ser His Leu Ala Leu Val Phe Val Lys Cys Phe
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 Ile Ile Gly Phe Ile Ile Leu Ala Ala Ile Leu Gly Asn Met Leu Val

115	120	125
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Phe Val Val Ser Leu Ala Val Ala Asp Met Leu Val Ala Leu Cys Ala		
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Met Thr Phe Asn Ala Ser Val Met Ile Ser Gly Lys Trp Met Phe Gly		
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Ser Val Met Cys Asp Met Trp Asn Ser Phe Asp Val Tyr Phe Ser Thr		
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Ala Ser Ile Met His Leu Cys Cys Ile Ser Val Asp Arg Tyr Tyr Ala		
195	200	205
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210	215	220
Phe Ile Met Leu Leu Met Val Trp Leu Ser Pro Ala Leu Leu Ser Phe		
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Lys Thr Leu Lys Ser Leu Phe Pro Tyr Ala Phe Tyr Phe Cys Arg Arg		
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<210> 70

<211> 3651

<212> DNA

<213> Drosophila

<400> 70

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<210> 71

<211> 1272

<212> DNA

<213> Drosophila

<400> 71

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<210> 72

<211> 423

<212> PRT

<213> Drosophila

<400> 72

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  35              40              45
His Tyr Ala Gly Ser Ser Asp Lys Leu Val Leu Leu Asp Asp Gly Arg
  50              55              60
Leu Arg His Tyr Thr Asn Ala Glu Asn Glu Ala Glu Glu Arg His Gly
  65              70              75              80
Ile Gln Ser Asp Tyr Glu Glu Asp Ile Ala Gly Ser Leu Glu Pro Leu
  85              90              95
Tyr His Asp Tyr Asp Lys Gly Leu Tyr Cys Ile Asp Lys Ala Thr Ser
  100             105             110
Ser Thr Gly Glu Glu Asn Val Leu Phe Ala Asn Ile Cys Leu Ala Arg
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Lys Glu Ile Lys Trp Ser Asp Ser Asn Phe Leu Leu Arg Lys Ile Leu
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  165             170             175
Ile Val Thr Thr Ile Ala Met Cys Leu Met Val Ser Gln Ala Ala Asp
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Leu Val Arg Ile Phe Thr Glu Leu Thr Ser His Val Ser Phe Ile Val
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<210> 73

<211> 5299

<212> DNA

<213> Drosophila

<400> 73

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<213> Drosophila

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<212> DNA

<213> Drosophila

<400> 80

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<211> 451

<212> PRT

<213> Drosophila

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<211> 7448

<212> DNA

<213> Drosophila

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<213> Drosophila

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<211> 6238

<212> DNA

<213> Drosophila

<400> 88

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<212> PRT

<213> Drosophila

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<211> 1019

<212> PRT

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Thr Asp Trp Ser Leu Arg Arg Glu Arg Asp Lys Leu Cys Ser Phe Ile
          35           40           45
Phe Lys Arg Val Leu Thr His Thr Glu Cys Lys Arg Leu Leu Glu Ala
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Ala Lys Ile Tyr Gly Glu Ser Ser Asp Phe Asn Ser Trp Pro Val Asn
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Glu Gln Leu Gln Trp Tyr Arg Asn Phe Asp Val Asn Asn Thr Asn Leu
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Gly Ser Ser Ser Asp Ser Glu Ile Leu Gly Pro Val Leu Pro Pro Phe
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Asp Arg Asn Ile Arg Gln Pro Ser Asn Tyr Phe Ile Ala Ser Leu Ala
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Trp Leu Ser Val Asp Tyr Thr Val Cys Leu Val Ser Gln Tyr Thr Val
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Gln Phe Leu Lys Asp Pro Ile Phe Asn Thr Ala Leu Ile Ile Gly Tyr
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Lys Thr Ala Tyr Asp Met Gln Lys Arg Ser Glu Ala Lys Gln Arg Lys
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Thr Val Ile Lys Arg Leu Ser Gly Ser Gly Gln Ala Asn Pro Leu Ala
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Gln	Ala	Ala	Leu	Leu	Ala	Thr	Arg	Gly	Lys	Gly	Asn	Leu	Gln	Lys	Ser
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Thr	Thr	Asn	Ser	Lys	Ser	Ile	Glu	Ala	Met	His	Gln	Tyr	His	His	His
			500					505					510		
Gln	Gln	His	His	His	Asn	Gln	Ser	Pro	Leu	Gln	Arg	Ala	Gln	Ser	Lys
		515					520					525			
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Ser	Gly	Asn	Ser	Glu	Leu	Ala	Leu	Thr	Tyr	Asp	Leu	Met	Thr	Asn	Ser
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 <213> Drosophila

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<211> 1014

<212> DNA

<213> Drosophila

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<210> 96

<211> 337

<212> PRT

<213> Drosophila

<400> 96

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 35      40      45
His Val Ala Arg Trp Asn Ala Thr Gly Asn Ala Thr Ile Ser Ala Thr
 50      55      60
Phe Glu Asp Val Pro Phe Asp Ala Asn Asn Tyr Trp Ala Leu Leu Ala
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Cys Tyr Ile Lys Cys Glu Lys Tyr Cys Ser Gly Ser Asp Arg Ser Ser
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Cys Cys Cys Ser Cys Thr Gly Thr Arg Pro Pro Thr Gly Gly Lys Lys
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<211> 9268

<212> DNA

<213> Drosophila

<400> 97

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<212> PRT

<213> Drosophila

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<211> 3731

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<213> Drosophila

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<211> 1167

<212> DNA

<213> Drosophila

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Val	Trp	Leu	Asn	Ala	Ser	Thr	Leu	Leu	Thr	Ser	Thr	Thr	Thr	Ala	Ala	225	230	235	240
Pro	Pro	Thr	Pro	Ser	Pro	Val	Val	Arg	Asn	Val	Thr	Val	Tyr	Arg	Leu	245	250	255	
Tyr	His	Ser	Asp	Leu	Ala	Leu	His	Asn	Ala	Ser	Leu	Gln	Asn	Ala	Thr	260	265	270	
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Gly	Thr	Lys	Ser	Pro	Ala	Asn	Gly	Lys	Ala	Ala	Asp	Arg	Pro	Arg	Lys	325	330	335	
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Met	Leu	Leu	Ala	Val	Leu	Leu	Leu	Phe	Leu	Ile	Thr	Glu	Phe	Pro	Gln	355	360	365	
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Gln	Cys	Tyr	Leu	Arg	Leu	Ser	Asp	Leu	Met	Asp	Ile	Leu	Ala	Leu	Ile	385	390	395	400
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Arg	Thr	Thr	Phe	Thr	Leu	Leu	Phe	Arg	Pro	Lys	Phe	Leu	Asp	Lys	Trp	420	425	430	
Leu	Pro	Val	Ala	Gln	Asp	Glu	Met	Ala	Ala	Ala	Arg	Ala	Glu	Arg	Ser	435	440	445	

Ala Val Ala Pro Val Leu Glu Lys Gly Arg Gln Gln Pro Gln Val His
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<211> 5251

<212> DNA

<213> Drosophila

<400> 109

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<211> 2325

<212> DNA

<213> Drosophila

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<211> 774

<212> PRT

<213> Drosophila

<400> 111

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100          105          110
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<211> 4314

<212> DNA

<213> Drosophila

<400> 112

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5927

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<211> 2325

<212> DNA

<213> Drosophila

<400> 122

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<210> 123

<211> 774

<212> PRT

<213> Drosophila

<400> 123

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35     40     45
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50     55     60
Glu Arg Ala Thr Leu His Asp Val Cys Leu Leu Arg Asn Gly Met Pro
65     70     75     80
Val Thr Arg Glu Cys Arg Val Lys Asp Asn Arg Ala Asn Trp Glu Ser
85     90     95
Thr Glu His Trp Asp Pro Val Val Cys Leu Arg Arg Phe Arg Glu His
100    105    110
Thr Ile Ser Gly Asp Leu Asn Ser Leu His Asp Asp Val Leu Glu Gly
115    120    125
Arg Arg Arg Thr Asn Asp Thr Gln Gly Arg Arg Glu Met Thr Gly Ile
130    135    140
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Val His Met Thr Gly Gln Met Phe Gly Ala Leu Met Gln Gln Asp Lys
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675	680	685
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<211> 7986

<212> DNA

<213> Drosophila

<400> 124

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<210> 125

<211> 1704

<212> DNA

<213> Drosophila

<400> 125

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<210> 126

<211> 567

<212> PRT

<213> Drosophila

<400> 126

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Gly Thr Leu Leu Leu Ile Gln Thr Ser Ala Ala Pro Gly Gly Asp Val
          35          40          45
Met Ala Thr Gly Ala Gly Ala Gly Lys Leu Gly Val Pro Ser Met Asn
          50          55          60
Ser Tyr Leu Val Ser Met Ala Trp Pro Lys Ser Leu Ala Val Ala Val
          65          70          75          80
Phe Leu Val Ile Ile Leu Val Thr Val Val Gly Asn Thr Leu Val Ile
          85          90          95
Leu Ala Val Leu Thr Thr Arg Arg Leu Arg Thr Val Thr Asn Cys Phe
          100          105          110
Val Met Asn Leu Ala Ile Thr Asp Trp Leu Val Gly Thr Cys Val Met
          115          120          125
Pro Pro Ser Val Val Leu Tyr Ile Thr Gly Thr Trp Arg Phe Gly Trp
          130          135          140
Ile Leu Cys Asp Ile Trp Ile Ser Leu Asp Ile Leu Leu Cys Ser Gly
          145          150          155          160
Ser Ile Leu Ser Leu Cys Ala Ile Ser Leu Asp Arg Tyr Leu Ala Val
          165          170          175
Thr Gln Pro Leu Thr Tyr Ser Lys Lys Arg Arg Ser Lys Arg Leu Ala
          180          185          190
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Pro Pro Tyr Leu Gly Trp Tyr Glu Val Gly Arg His Gln Ala Glu Phe
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Asn Arg Lys Ser Arg Trp Arg Phe Asn Ser Leu His Asp Pro Pro Pro
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 Thr Pro Ser Gly Met His Tyr Val Gly Thr Glu Ala Ser Phe Ser Asp
 405 410 415
 Thr Cys Leu Gly Gly Asn Lys Leu Glu Ala Glu Ala Lys Gln Lys Gln
 420 425 430
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 His Gly Ser His Gly Gln Gln Ser His His Asn His His His Arg Met
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 Val Tyr Tyr Leu Leu Ile Pro Phe Leu Pro Arg Pro Ala Val Leu Glu
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 Asp Leu Met Phe Gly Phe Thr Trp Ile Gly Trp Val Asn Cys Ala Ile
 515 520 525
 Asn Pro Phe Ile Tyr Ala Phe Tyr Asn Pro Asp Phe Arg Thr Ala Phe
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<210> 127

<211> 3681

<212> DNA

<213> Drosophila

<400> 127

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<210> 128

<211> 1503

<212> DNA

<213> Drosophila

<400> 128

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<210> 129

<211> 172

<212> PRT

<213> Drosophila

<400> 129

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      20             25             30
Ile Ser His Ile Pro Lys Leu Asn Asp Ser Tyr Ala Tyr Glu Glu Leu
      35             40             45
Ile Ile Pro Ala His Leu Thr Gly Leu Tyr Thr Phe Arg Gln Leu Ala
      50             55             60
Asp Gly Ser Gln Glu Pro Val Lys Ser His Leu Arg Ala Cys Ile Cys
      65             70             75             80
Lys Leu Lys Pro Cys Ile Arg Phe Cys Cys Pro Arg Asn Lys Met Met
      85             90             95
Pro Asn Ser Arg Cys Ser Asp Gly Leu Thr Glu Asn Leu Lys Arg Ile
      100            105            110
Asn Pro Tyr Leu Lys Ile Thr Leu Glu Asp Gly Thr Ile Gly Lys Tyr
      115            120            125
Tyr Leu Leu Thr Asp Met Ile Val Leu Arg Tyr Glu Phe Arg Tyr Cys
      130            135            140
Glu Lys Val Val Ser Val Gln Glu Asp Gln Tyr Lys Leu Tyr Glu Val
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Lys Ile Ser Ile Phe Leu Lys Tyr Leu Ser Ile Ile
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<210> 130

<211> 6552

<212> DNA

<213> Drosophila

<400> 130

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<211> 1392

<212> DNA

<213> Drosophila

<400> 131

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<210> 132

<211> 444

<212> PRT

<213> Drosophila

<400> 132

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His Lys Asn Ser Arg Arg Val Cys Leu Thr Ile Leu Leu Val Trp Ala
 35           40           45
Ile Ser Ala Ala Ile Gly Ser Pro Ile Val Leu Gly Leu Asn Asn Thr
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Pro Asn Arg Glu Pro Asp Val Cys Ala Phe Tyr Asn Ala Asp Phe Ile
 65           70           75           80
Leu Tyr Ser Ser Leu Ser Ser Phe Tyr Ile Pro Cys Ile Ile Met Val
 85           90           95
Phe Leu Tyr Trp Asn Ile Phe Lys Ala Leu Arg Ser Arg Ala Arg Lys
100           105           110
Gln Arg Ala Ala Arg Lys Pro His Leu Ser Glu Leu Thr Gly Gly Ser
115           120           125
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Asn Thr Ala Ser Gly Ser Asn Glu Glu Glu Asp Glu Asn Ala Ile Ser
165           170           175
Pro Asp Ile Asp Asp Cys His Val Ile Val Asn Asp Lys Ser Thr Glu
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Phe Met Leu Ala Thr Val Val Glu Glu Thr Gly Asn Ser Val Val Ala
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Gln Ile Thr Thr Gln Pro Gln Leu Val Val Ala Asp Pro Asn Gly Asn
210           215           220
His Asp Ser Gly Tyr Ala Ala Ser Asn Val Asp Asp Val Leu Ala Gly
225           230           235           240
Val Ala Pro Ala Ser Ala Ser Ala Ala Thr Ser Ala Ala Pro Arg Ser
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Ser Gly Ser Pro Pro Asp Ser Pro Leu Pro Ser Gly Ala Thr Leu Gln
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Arg Ser Ser Val Ser Ser Gln Arg Arg Pro Thr Gly Asp Asp Ser Pro
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Lys Arg Gly Glu Pro Ala Leu Ser Val Ala Met Lys Pro Leu Ser Phe
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Val Arg Tyr Gly Val Gln Glu Ala Met Thr Leu Ala Arg Asn Asp Ser
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 Thr Leu Ser Thr Thr Ser Lys Thr Ser Ser Arg Lys Asp Lys Lys Asn
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<210> 133

<211> 2970

<212> DNA

<213> Drosophila

<400> 133

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<210> 134

<211> 1116

<212> DNA

<213> Drosophila

<400> 134

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<210> 135

<211> 371

<212> PRT

<213> Drosophila

<400> 135

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Thr Glu Thr Pro Thr Glu Ile Thr Leu Gln Ala Thr Ser Phe Gly Ala
 35          40          45
Gly His Leu Leu Trp Leu Ala Ile Asn Ala Phe Leu Phe Val Leu Ile
 50          55          60
Leu Gly Gly Asn Ile Leu Thr Ile Val Ala Val Arg Thr Cys Arg His
 65          70          75          80
Leu Arg Ser Val Ile Ser Asn Leu Phe Ile Leu Ser Leu Ala Val Ser

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<211> 1365

<212> DNA

<213> Drosophila

<400> 137

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<211> 444

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<213> Drosophila

<400> 138

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210    215    220

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<211> 4185

<212> DNA

<213> Drosophila

<400> 139

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<211> 1584

<212> DNA

<213> Drosophila

<400> 140

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<210> 141

<211> 322

<212> PRT

<213> Drosophila

<400> 141

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Gly Leu Leu Val Val Pro Phe Ser Val Tyr Pro Ala Leu Thr Gly Glu
65     70     75     80
Trp Met Tyr Gly Asp Ile Val Cys Arg Phe Thr Gly Tyr Leu Glu Val
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Ser Ala Thr Leu Ala Ile Leu Val Leu Gly Pro Ser Leu Ile Ser Ile
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195    200    205
Gly Glu Pro Ile His Asp Lys Glu Tyr Ala Thr Ala Leu Ala Glu Asn
210    215    220
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<210> 142

<211> 4900

<212> DNA

<213> Drosophila

<400> 142

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<211> 1245

<212> DNA

<213> Drosophila

<400> 143

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<210> 144

<211> 414

<212> PRT

<213> Drosophila

<400> 144

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<211> 4116

<212> DNA

<213> Drosophila

<400> 145

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 <213> Drosophila

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<210> 153

<211> 706

<212> PRT

<213> Drosophila

<400> 153

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  20          25          30
Ile Gly Ile Ala Glu Thr Ile Glu Ala Val Leu Ile Leu Val Leu Thr
  35          40          45
Leu Gly Val Ile Gly Ala Asn Cys Leu Val Ile Phe Val Ile Asn Asn
  50          55          60
Arg Arg Tyr Ala Ala Tyr Ile His Gln Gln Pro Arg Tyr Leu Leu Thr
  65          70          75          80
Ser Leu Ala Leu Asn Asp Leu Thr Ile Gly Leu Leu Ile Thr Pro Phe
  85          90          95
Gly Leu Met Pro Ala Leu Phe His Cys Trp Pro Tyr Gly Glu Ile Phe
 100          105          110
Cys Gln Ile Gln Ala Leu Leu Arg Gly Ala Leu Ser Gln Gln Ser Ala
 115          120          125
Val Ile Leu Val Cys Met Ala Val Asp Arg Tyr Met Cys Ala Leu His
 130          135          140
Pro Arg Arg Tyr Tyr Gln His Ser Ser Lys Lys Gly Cys Val Ala Ile
 145          150          155          160
Leu Ser Leu Thr Trp Ile Ile Ser Leu Thr Val Phe Gly Phe Leu Val
 165          170          175
Leu Pro Lys Gly Tyr Tyr Phe Asn Asn Thr Gly Leu Met Ala Cys Glu
 180          185          190
Pro Phe Tyr Ser Lys Pro Ser Tyr Arg Ile Leu Ser Thr Cys Ala Leu
 195          200          205
Tyr Phe Pro Thr Thr Met Val Leu Met Tyr Cys Tyr Gly Ser Ser Phe
 210          215          220
His Met Ser Arg Phe Arg Leu Asn Asp Pro Thr Met Pro Leu Thr Ala
 225          230          235          240

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Ala Ala His His Pro His Pro His Pro His Pro Thr Ala Ala Gln Gln
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 405 410 415
 Tyr Trp Leu Leu Asn Ser Asp Phe Arg Arg Met Ser Arg Gln Leu Met
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 Cys Cys His Ile Asn Ala Asn Asp Phe Glu Ile Thr Thr Leu Pro Ile
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 Asn Ser Leu Ser Arg Ser Ala Ser Gln Tyr Ile Arg Gly Thr Met Gly
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 545 550 555 560
 Arg Thr Val Ser Ser Gly Asn Leu Asn Ala Met Gln Lys His Leu Pro
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 595 600 605
 Leu Gln Leu Asn Leu Asn Leu Ser Gln Ala Ala Thr Ala Ala Glu Leu
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 Gly Lys Phe Ser Asn Ser Glu Pro Lys Leu Cys Glu His Leu Phe His
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 Asp Tyr Ala Glu Asp Val Ile Leu Ala Lys Asn Gln Leu Leu Ala Arg
 645 650 655
 Gln Ala Lys Cys Ser His His Pro Leu His Gln Gln Ala Lys Thr Arg
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<210> 154
 <211> 3332
 <212> DNA
 <213> Drosophila

<400> 154

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<210> 155

<211> 1332

<212> DNA

<213> Drosophila

<400> 155

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<210> 156

<211> 393

<212> PRT

<213> Drosophila

<400> 156

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 20          25          30
Ser His Arg Asn Asn Ser Thr Arg Thr Asn Ile Ala Thr Asn Gly Cys
 35          40          45
Ala His Ser Gly Ile Leu Leu Phe Val Leu Thr Ala Met Thr Leu Thr
 50          55          60
Ser Leu Ile Thr Pro Thr Glu Gln Leu Ala Val Ala Pro Asn Gly Thr
 65          70          75          80
Thr Leu His Gln Leu Glu Ser Val Glu Ser Glu Ser Tyr Pro Ser Ile
 85          90          95
Asn Gly Thr Gln Asn Glu Thr Met Val Thr Ser Val Arg Pro His Leu
100          105          110
Asp His Arg Asn Arg Pro Thr Gln Gln Asn Gly Ser His Tyr Leu Glu
115          120          125
Tyr Asp Asp Asp Gly Pro Asp Cys Ser Tyr Ser Tyr Asn Phe Ile Leu
130          135          140
Lys Leu Ile Thr Met Ile Leu Tyr Ala Leu Val Cys Ile Ile Gly Leu

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Phe Gly Asn Thr	Leu Val Ile Tyr	Val Val Met Arg	Phe Ser Lys Met			
	165	170	175			
Gln Thr Val Thr	Asn Ile Tyr Ile	Leu Asn Leu Ala	Ile Ala Asp Glu			
	180	185	190			
Cys Phe Leu Ile	Gly Ile Pro Phe	Leu Leu Tyr Thr	Met Gln Val Gly			
	195	200	205			
Asn Trp Pro Phe	Gly Asn Tyr Met	Cys Lys Ala Tyr	Met Val Ser Thr			
	210	215	220			
Ser Ile Thr Ser	Phe Thr Ser Ser	Ile Phe Leu Ile	Met Ser Ala			
225	230	235	240			
Asp Arg Tyr Ile	Ala Val Cys His	Pro Ile Ser Ser	Pro Arg Tyr Arg			
	245	250	255			
Thr Pro Phe Val	Ser Lys Leu Val	Ser Ala Phe Ala	Trp Met Thr Ser			
	260	265	270			
Val Leu Leu Met	Leu Pro Val Ile	Leu Phe Ala Ser	Thr Val Gln Ser			
	275	280	285			
Ser Asn Gly Asn	Val Ser Cys Asn	Ile Glu Trp Pro	Asp Thr Gln Asn			
	290	295	300			
Ser His Thr Asp	Ser Thr Phe Ile	Leu Tyr Ser Leu	Val Leu Gly Phe			
305	310	315	320			
Ala Thr Pro Leu	Thr Phe Ile Leu	Val Phe Tyr Cys	Leu Val Ile Arg			
	325	330	335			
Lys Leu His Thr	Val Gly Pro Lys	His Lys Ser Lys	Glu Lys Lys Arg			
	340	345	350			
Ser His Arg Lys	Val Thr Lys Leu	Val Leu Thr Val	Ile Ile Tyr Asp			
	355	360	365			
Ile Ser Ile Ser	Cys Gly Thr Asn	Tyr Tyr Leu Thr	Thr Leu Val Phe			
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<210> 157

<211> 5453

<212> DNA

<213> Drosophila

<400> 157

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<211> 1902

<212> DNA

<213> Drosophila

<400> 158

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<213> Drosophila

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<211> 4385

<212> DNA

<213> Drosophila

<400> 160

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<212> DNA

<213> Drosophila

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<210> 162

<211> 462

<212> PRT

<213> Drosophila

<400> 162

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Val Cys Ile Leu Gly Thr Ile Ala Asn Thr Leu Asn Ile Ile Val Leu
35          40          45
Thr Arg Arg Glu Met Arg Ser Pro Thr Asn Ala Ile Leu Thr Gly Leu
50          55          60
Ala Val Ala Asp Leu Ala Val Met Leu Glu Tyr Ile Pro Tyr Thr Val
65          70          75          80
His Asp Tyr Ile Leu Ser Val Arg Leu Pro Arg Glu Glu Gln Leu Ser
85          90          95
Tyr Ser Trp Ala Cys Phe Ile Lys Phe His Ser Val Phe Pro Gln Val
100         105         110
Leu His Thr Ile Ser Ile Trp Leu Thr Val Thr Leu Ala Val Trp Arg
115         120         125
Tyr Ile Ala Val Ser Tyr Pro Gln Arg Asn Arg Ile Trp Cys Gly Met
130         135         140
Arg Thr Thr Leu Ile Thr Ile Ala Thr Ala Tyr Val Val Cys Val Leu
145         150         155         160
Val Val Ser Pro Trp Leu Tyr Leu Val Thr Ala Ile Ala Lys Phe Leu
165         170         175
Glu Thr Leu Asp Ala Asn Gly Lys Thr Ile Ala Ser Val Pro Leu Ser
180         185         190
Gln Tyr Ile Leu Asp Tyr Asn Arg Gln Asp Glu Val Thr Met Gln Val
195         200         205
Met Ser Ser Thr Thr Pro Asp Val Ser Trp Ala Ile Pro Ile Thr Thr
210         215         220
Ser Ser Ser Leu Gly Glu Arg Asn Val Thr Val Tyr Lys Leu Tyr His
225         230         235         240
Ser Ala Leu Ala Leu Arg Asp Arg Gln Phe Arg Asn Ala Thr Phe Leu
245         250         255
Ile Tyr Ser Val Leu Ile Lys Leu Ile Pro Cys Phe Ala Leu Thr Ile
260         265         270
Leu Ser Val Arg Leu Ile Gly Ala Leu Leu Glu Ala Lys Arg Arg Arg
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Lys Ile Leu Ala Cys His Ala Ala Asn Asp Met Gln Pro Ile Val Asn
290         295         300
Gly Lys Val Val Ile Pro Thr Gln Pro Lys Ser Cys Lys Leu Leu Glu
305         310         315         320
Lys Glu Lys Gln Thr Asp Arg Thr Thr Arg Met Leu Leu Ala Val Leu
325         330         335

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 Asp Gly Glu Gly Arg Val Gly Gly Ser Gly Gly Leu Gly Gly Tyr Gly
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<210> 163

<211> 4214

<212> DNA

<213> Drosophila

<400> 163

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4214

<210> 164

<211> 1859

<212> DNA

<213> Drosophila

<400> 164

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<210> 165

<211> 513

<212> PRT

<213> Drosophila

<400> 165

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20     25     30
His Thr Val Asn Ile Thr Asp Gly Leu Arg Met Lys Asp Gly Ser Tyr
35     40     45
Ser Tyr Ala Gly Val Val Val Pro Pro His Leu Met Ala Glu Tyr Ser
50     55     60
Phe Lys Val Ile Asp Gly Val Glu Tyr Arg Ala Lys Lys His Leu Arg
65     70     75     80
Gly Cys Val Cys Leu Leu Lys Pro Cys Ile Ser Phe Cys Cys Pro Glu
85     90     95
Asn Leu Val Phe Asp Ala Lys His Trp Asn Cys Thr Met Pro His Gln
100    105    110
Val Arg Glu Ser Thr His Val Glu Leu Thr Tyr Ala Asn Arg Thr Val
115    120    125
Asp Gln Val Arg Ile Arg Asp Arg Phe Val Val Arg Thr Glu Leu Gly
130    135    140
Cys Arg Asn Lys Phe Val Asp Lys Lys His Asp Asn Phe Trp Gln Trp
145    150    155    160
Asp Leu Phe Glu Asn Gly Thr Leu Arg Arg Asp Asn Arg Leu Trp Ser
165    170    175
Thr Asp Glu Tyr Cys Phe Ser Pro Leu Glu His Asn Pro Glu Gln Trp
180    185    190
Glu Leu Thr Pro Leu Asn Cys Glu Arg Phe Gln Thr Gly Tyr Arg Val
195    200    205
Trp Ile Tyr Ala Ile Cys Ser Ile Ile Ala Ile Ile Ile Asn Ile Phe
210    215    220
Ile Leu Ser Leu Leu Gly Ser Val Arg Asp Ala Arg Lys Ser His Tyr
225    230    235    240
Gly Gln Leu Ile Ile Tyr Tyr Leu Leu Ser Met Ile Val Gly Tyr Ser
245    250    255
Leu Leu Val Tyr Leu Ala Leu Lys Asn Pro Met Lys Leu Ser His Val
260    265    270
Ala Cys Arg Asn Ile Gly Phe Leu Ala Tyr Phe Cys Ile Met Leu Ser
275    280    285
Phe Val Phe Leu Ala Ile Cys Ser Leu Asp Phe Leu Leu Lys Phe Lys
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Gln Lys Ala Val Arg Ser Ser Val Arg Arg Leu Ser Leu Ala Leu Ala
305    310    315    320

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 Phe Ser Ile Tyr Glu Leu Pro Pro Asp Thr Gln Tyr Ile Leu Gly Thr
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 Gln Leu Lys Ile Val Lys Thr His Phe Tyr Ala Phe Ser Ala Tyr Ile
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 Val Gly Val Phe Ala Val Trp Ile Arg Glu Ile Val Val Tyr Ile Met
 420 425 430
 Ala Arg Val Arg Glu His Phe Phe Ile Ile Asp Phe Trp Ser Gly Ile
 435 440 445
 Cys Ile Leu Gly Leu Ala Ile Ala Gly Phe Ile Leu Leu Leu Gly Lys
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 Asn Leu His Val Lys Ser Trp Trp Ala Ile Asn Val Glu Ser Ser Gln
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<210> 166

<211> 5208

<212> DNA

<213> Drosophila

<400> 166

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5208

<210> 167

<211> 1914

<212> DNA

<213> Drosophila

<400> 167

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<210> 168

<211> 637

<212> PRT

<213> Drosophila

<400> 168

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 35           40           45
Ile Asp Asp Pro Asn Val Pro Cys Asn Phe Tyr Asp Thr Val Asn Leu
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Thr Gly His Arg Leu Phe Pro Asn Gly Ser Tyr Asp Tyr Tyr Gly Thr
 65           70           75           80
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213

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<211> 3957

<212> DNA

<213> Drosophila

<400> 169

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<210> 170

<211> 1638

<212> DNA

<213> Drosophila

<400> 170

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<210> 171

<211> 545

<212> PRT

<213> Drosophila

<400> 171

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 35      40      45
Gly Asn Phe Ser Leu Gly Asn Pro Tyr Asp Val Asp Ser Glu His Ser
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Arg Arg Lys Leu Cys Ala Leu Ser Leu Thr Ala Arg Ala Ala His Gly
 275     280     285
Glu Leu Pro Leu Pro Ile Pro Ile Leu Arg Arg Gln Thr His Met Val
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Ser Ser Thr Asn Glu Ala Ala Leu Gly Ala Phe	Asn Pro Lys Leu Ile	
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<210> 172

<211> 7112

<212> DNA

<213> Drosophila

<400> 172

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 Gln Ala Lys Gly Ala Ala Lys Lys Lys Arg Leu Met Lys Gln Leu

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SUBSTITUTE SHEET (RULE 26)

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<210> 182

<211> 2010

<212> DNA

<213> Drosophila

<400> 182

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<210> 183

<211> 669

<212> PRT

<213> Drosophila

<400> 183

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Gly Lys Phe Pro Asn Ala Asn Thr Pro Ile Ala Ile Asp Glu Pro Thr

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SUBSTITUTE SHEET (RULE 26)

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<210> 184

<211> 4243

<212> DNA

<213> Drosophila

<400> 184

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<210> 185

<211> 1224

<212> DNA

<213> Drosophila

<400> 185

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<211> 407

<212> PRT

<213> Drosophila

<400> 186

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		20					25						30		
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Val	Ile	Gln	Asn	Pro	Thr	Gln	Cys	Ser	Phe	Val	Val	Asn	Lys	Tyr	Tyr
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 <211> 11847
 <212> DNA
 <213> Drosophila

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<211> 6617

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<213> Drosophila

<400> 190

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<211> 1983

<212> DNA

<213> Drosophila

<400> 191

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<211> 4902

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Ser Ser Val
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Exon: 2674..2584
Exon: 2518..2427
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2/89

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Name: Calcitonin receptor-like
Classification: G_protein_linked_receptor

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FIGURE SHEET 3

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Exon: 4058..4202
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Exon: 7120..7228
Exon: 7333..7504
Exon: 7648..7754
Exon: 8263..8412
Exon: 8583..8716
Exon: 8953..9143
Exon: 9801..9935
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Exon: 10294..10507
Exon: 10885..11036
Exon: 13701..13884
Exon: 14430..14572
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(SEQ ID NO: 6)

Name: metabotropic glutamate receptor-like
Classification: G protein linked receptor

GenBank sequence NO.: U42800.1

CGATAGTTCTCTAGTTACCAAGTGGAGTTTGGTTGAAATCTTCTTTGCTATATAGGTATTTTCATTTCCGATATGCTGTATCGGTTTGTAAAATAATAAAGCATTTCCGCAGCTCTCTCTCTCTCTGCTATCTCAAAATAATAAGTAATGCCCATATGTGTGGTCTTAAGCATCTAGATTGATGAAGAAATGACACTTCAATCGCTCCGCTCTGTTTGTCTTATCAACGCTAGCTGAATTCAGCAACAACAATGGCAATGTATTATTTCCCTTTACTTCTTCTTATCTCTTATTGATTCGCGATTTGGAAGAAATTTAGTTGTATAGTGGAAAAACGGCAGCTGGATAGCAAGAAATGACAATCTTTCAAACGATAGGATATCTTTTGGTTTATGAACAACAAAACAAATCTTAAGCCTGTGTGTATATATATATATATCTTACGAGCATTAATCAGCGTAAATCAAAAACAAATGATTTCTTCAAAAACATAAAACATAATTCGCCGTGAATCTTATATATCATATGATACACAGTTCCTAGATCTTATACATATACCTTACATCCCTTCCATATATATATATCTTGATTTCTGTTTGTAAATAGTTTTTTAAAGTATTTTAAAGTATTTAAATAGGCATGTGGTGATGTATTTACAATACATACATACATATATCGTACATACATCTTATTTAATATCGATTTCCAAGTTGGGTATAAAATGCATGCAAGTCGAAGGGCCCGATTCCAATTCCAATTCCAACCCCCCATCGCTCCAGATTTTGGTTAAGCTTGCACGGCATCCGAGGGGTATGGATGTCCCATCTAGGTCTAGTTTGTATTGATTTCATATACATTTTAAAGTTTGGGATTTTTCGAATCTCGCAGTATAGCATAGCATGACAGATGTTGTTGTTACACATATGCTTCATATGCGCTCAACGACTAATAC TAGTATAGCTAGAGATCCTAAGCTACAATTTTGCCTAAGACTGCTCCGGGGGAATTCGCTTTACTGATCACTTAG

[illegible]

Start ATG: 1 (Reverse strand: CAT)

MLPTILISYEHISLDLSKYQTAYACEGKKLTIECDPGDVINLIRANYGRFSITICNDHGNVSVNCMFPKSLSVLNSRCAHKQSCGVLAATSMFEGDPCPGTHKYLEAHYQ
CISAAQTSTTNRPSPPWVLSNGPPIFGNGSLIHPPGVGAGAPPPRLPTLPGVVGISGNPGLFNVPQHTATVHTSTPSSSTTAVGGGRLLKGGATSTTTKHPAGRHGDL
PPPQLHHHHHHHGEDTASPTKSSKLPAAGNATSPSNTRIILTVGGSGSDTGGTLTTKSSPNRPGTAASGSVVPGNGSVVRTINNINLNAAGMSGGDDSEKLFCCPGTHAR
NLYNMNTRVGDVNVQPCPGGAAGIAKWRLMKRIPDSGYDEYDDDISSTTPAPSGGDCLNHSSSCEPVMSAHKVNMQRRLNFEPTHMNPAFDPLTQCRSLWNLEMRMVRN
DSSLISITANDMSEPTSSKTLVGGDMLVTTKIIQTVEKMMHDKETEPDQOREAMIMELLHCVVCTGSNLLDSEQLSWLDLNPEDQMRVATSLTLTGLEYNAFFLADTIRE

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(SEO ID NO: 9)

Name: Latrophilin receptor-like
Classification: G protein linked receptor

(SEQ ID NO: 10)

Exon: 3914..3576
Exon: 3322..2232
Exon: 2170..1932
Exon: 1854..1001
Start ATG: 3241 (Reverse strand: CAT)

transcript: chr2: 119500
GACCGATTCGGTGAATTTTACCACGAAAAACAAATAAAATATATAACCTTCGATGTTAATACCAATTAGATTAGAATACATTCGCTAAGGCTACAACTGACTGCGCCA
TCGGTTTAAGCACATCGAAACAAATACACTATTCGTCTACAGATGTTGTGAAGAAGTTCGATGTTGCTGATAGCTGTTTCGGGTTAATCCCTGAGGAAACCAACCGCTTATC
GACGTCGATCAGCGGGTTGGCGCGCGCTCGGGTGAAGACAGGGGAGTGGTTGGGGCTGAAGAAGCTGTGGGATGAGGAGGAGGAGGAGCAACAGGACATAACAGACACGCGCGAAAAA

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GAGAGAAACGGAATCGGAACACCAAGCGAAGAAACGTATCGAATTCGTATTCTTGGAAACGAAAGAACAAACAAATCAGCCGCAATGAAGCCAACCTGCATATTGTGTCTCC
 TGGTGGTGATCCTCCTGCATCCGCGCATCTCAAAATCTTCAACGAGCGGCAACCCAGTGCCAGTTCAGTTCAGTTCCTCCGCCAGAAATCCCGCCTTCCGGCAATGCGA
 AACCATTCGATTGAGATGTGCCGCAAGATCGGCTACAATGAGACCTCCATGCCAAATCTGGTGGGTAACGAAATGCAACGAGCTGGAGTACACACTGCAAACTTTGCA
 CCACTGATCGAATACGACTGCGATTCCAGTTGAAGCTGTTCTGTGCGCGCCTACGTGCCATGTGCACGCCAAAGCTCCAGTCCATGCCATCGGTCCGTGCCGGAGTC
 TCTGTAGTCCGTGCGAATACGTTGTATCCAGTTTTCAGGGATTTCGGATTTCATGGCCACCGGCTCTCGACTGCGATAAATTTCCGCGAGAGAACAAACACGAGACGAT
 GTGTATGGAAGGTCCCGGTGAATTGCATCAGCCGAGCAGGAGCAGGATTGTATGGGCTACCGGGCCAAAGGAATACCCGGTGGATTGGGCGGTAACTGCCCATGGATTGC
 TCGGACTGGCCAAATCGCATCTGTACGTAAGCTACCCAGATCTGGACGCTGTGCACCACTCTGCGAGGCGGACATACTGTTACACCGCGCGGAGAACATCTGGCAGAGA
 TCTGGGTCTCCACATGGGCTATGCCGCTTTGGGTTGGCCCTGGTAGCCACCGTGTGCTTTTGGCCAGCGATGGCAGTGGTTGGCCAGCGCCAGTGGTCCAGATTGTT
 GTCGCCGCTGATTTGGTGCCCAACATGGTCACCTTGGGCTGGGCGAGTGGCTTTATGGTGGGAAGAACGGGAATGCGTGGGCGACGGATCCCCAGGCGCCCAATGAATCC
 CTACTCACCCTGGACGAGTGTCCATGCTCTCTGCGTAGTGTCTCTTATGCGCTACTATTTGGAATGGCCGCTGCGCTTGGTGGGCTGTGCTCTGTTTGGGCTGGC
 ACCGCGACATTCGTCGGCAGTCCGGACTCCAAAGGCCATGTGGTCACTTCCCTCGAACTTTGGAGGCGAGTCCGCGGAAGAGGAACAGTGCCAAAGACGGCTCAGCAGGATCT
 AACGCAAAACAACTTTGTGTGCTTCGTGGCCTGGGATTTGCCGCGCTTTTCAAACCTCAGCAGTGTATGTTGCCGCTTTCGTGGATGCTGACGAGTTGCTTGGCACTAGTCTG
 GATAAGGCGCTACAGATATTGGTGGCTACGCCGCTTTTGTGCTAGGATCTTTGGCTCGATGAATCTGATCTCCGCTATTTGGTACACTGTGCGACCAAGGAGATCTTGA
 GGAATGCAATGCACTCAGCTGCAACAGCAACTGCAACAGCTAAGCGGACACAGTAGTTCGGCATTTGGCATATTCTTTTATCTATGAGCATTTGATGCTGCTGCTCTCT
 GTTGGCTGCTATTTACGATTGTCACCATCGATGTCTGGCTGGGATCGGGTATACCAATACGCCACTTTTGGCCATTTTTGTGGCGCTTCTATGGAACGATGATGCTGGG
 ATTTGTGCTTTTGGCTGGGTACTTGGACCAAGCATTTCACCTCTCTACAAGAGGCAAGTCAGCAATGCAAAATGGTGAAGCATACTGCTGCGGAGCAGCAACTGGACACC
 TGGACGGGCTAGTACCTCGCTGGATCGCATGCGGCTTGCATAGCAGCGGTGCTCTTACCATTCCGTTGCGGAATCTTATGGCCAGTGTTCGCTTGGCGGCAAGTCCGTA
 CAACTGAAGACTTTCGCTGCTGATCGCTGAATCAGATCTCCAACTTTCGCTGGGCGAGGCGTGATCACCAGGAGCGTGATCACCAGCGGCTATCATCATCATCATCAT
 CACCAGCAGCAGCAT
 GCCAGCAGCGGCTACTATCCCCACTTGCAGCAGTATGGCAATGAGACGCTACTCTAG
 (SEQ ID NO: 11)

Start ATG: 421 (Reverse strand: CAT)

MKPTCILLLVILLHPRIKSSSTSGNPSASSSSSSPPEIPAFRCQETIRIEMCRKIGYNETSMPNLVGNEMQTDVEYTLQTFAPLIEYDCSSQLKFLCAAYVPMCTPKAP
 VHAIGPCRLCESVRIRCHPVLQGGFPPWPPALDCDKFPRENNHETMCEGPELHQPQQEQDLYGLPGQIPGGGLGKLFMDCSGLAKSHLYVLRPSGRCAPICEADILF
 TPAEKHLAAEIWSTWYAAALGLALVATVCLLASDGRSLASAKVRLSLPLIACHNMVTLGWAVRFMVGRGTGTCGDPQAPNESLLVLDGLSNASCASVFLMRYVFGMAACA
 WWAVLCLGWHRRDIRHSPDSKGVHVIPSNFGGSPAKRNSAKTAQOQLTQNFVCFVWGLPAFQTSAVIIVARFVDADELLGNQSDKALQILVATPVFCYWIYFGSMNLISGLY
 VHCRTKEILRNSNLSVQQQLQQLSAHSSSGIGIFLFIYGLACAMLLAVIYEFANIDVWLGSGDTNTPLPWFLRLAFMELMLGICCFWVGLPSISLTKYRQVNSNGKMKVH
 TSAGAAATGHLGHSRSGSHAACNSTVHSHSVRTSMASVPLPPSPYKLTSPGTSISLQMSNYSLGRSVHQQRHSPPHHHHQQQHHQHFPHNNHHHSTSSSHRLYYF
 PGSYASQKYSQHSYYPHLQYGNETLL*
 (SEQ ID NO: 12)

Name: Frizzled 4 -like

Classification: G_protein_linked_receptor

Celera Sequence No.: 142000013384825

TCATACTACTGGCTGCGCAACCCCTACTCTACTCTGACTACTTTGGTTCTGGTTTGGATCTTGGCTGATGCTAGGTTCTAACTTTAAGTACGATTCTTAGTATTGCTCACT
 GGATACACTGTAGATACACCGCAGATACAGCCGCTCTGCGGCGGCACAACAAGAAACAAACCGTAAATGCAGCTACTTACGACTAGGAGGATGATTAGTGCAAGGGGT
 TAAGGAGCGGGTTAATCACAACCTGGCTGGGAACTGAAAGTGGCGCACAAGCGCATCTACTTAACTAGATGATGGTCTACGCGGCGCCAGGGTCAATCAGATTGCGGAGTA
 GGTCTACACAGAGAGATGCGATTGGTCAAAAACCTAAGCTTAAACGACCCGCGATGTTATCTTATCAATTAACGAAACGAAACGAAATGGAATCAACATTTTATAGACCCATTTA
 AAGAGCTATCATCTTTTACCAGAACTAATTAAGAAATAATATGATGCTTAGTAAAGGAACACTACACTAGGTTCTAAGAAATGATTATCAAGAAATTAACATTGGCA
 CCCCAGGCGGAATATGATCTGGTAGTTTTAGAAAACCTTTGCTGATACCGAAACTCGCGAATTAAGATAAATTTCAACATTGCAACAATTTCTTAGTTTACATTGTAA
 TTTTGTCTACTTTAGGAGAGCACAATGTGCTGCAATTTCTCCGCTGTGTGTGATCTGCTAAGTTTCAAGAAATGACTAGCTATGTGAATCTGACTTGAAGAGGAGGG
 GGAGGTTGATCTGCTGACTTATCTAGCTTTTACCCTAATGCATTGATTGCTTAAAGCTGATGATTGGTCAGTTGCACACGCGCTATCTGGGCGAGATTGTTGGGCGGTG
 TCGTAAAAAATGCTTGGTGTTCGCTTCAATTTGACGAGGACTTGGACTTGGATTGGAAGTGTGGACCGTGTACAGAGTGGACAAATGGACAGGGCATCTTATTTATTCGG
 TGCCATTGCGAGGCGCTGCTGCTCTTCTCGGAAATCGCGAGGAAAGCCTGCACTTGCCTGCTTTCCATCGATTTCAGGTGCTCCACTGGTGGCGCTGCTGAG
 CGAAATGCGAGGTATGGAATTTGACGCTGCCCAAGATAGAGGTGCTATCTCCGGGAGCGGCTCTCGGATTGCGCCAGGCGCGTGAGACTGCAAGTGGGCGGCAGGATGCGGTG
 GGAAAGCGTGGCGAACCCTTACCCTTTCCCTTAACTGGGTGGCTGCTGCCACCAGGAGTCTCCGCTGCTGGCAATTTGAAGTTTCACTTGGTCTCCAGCAGTTTCA
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 CACCTCCACTCGATACTGCAGTTCTCGGCGATCATCGACCAGAAATCGGCGTGTGTGCACTCCATCTTAAGGCCCATACTGTGGATTAACCGGATCGATAATATCGTCTGAG
 GGAGAAGGAGAGTTCCCAAAGGATTAGGACACAACCTTTATAAAGGTTCTAGAAATAGGAGACTAGAATTAAGATATTTCTAAGTGAGCAAGATCTTATTGACCTTCA
 AGCGCTACTTACTTCTGCCCAAAATATCATGGAGTTTGGGATGAAAGCAGCAGCAAGGGTTGCACTGGTGTGTTAAATCAGAGCTGTGATTGTGATAGAGGCGAGGT
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 ATAGATAGATAGATAGATGATTCTAATGATTCAATGATTAAATGATTCTTCAACATTAATTAAGTAATCTGATAATTTGAAGCATATTAATATTTTCTTACCTGAGGCTG
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 AATCCGCGCAGGAGCAAGATCAGTTGAATGCTCTGCAGCATCTTGTCTTGAAGACGCTGCGGTTACGAGTGAATACTCGATGCACTCTGTAGGTTCTTGGCAACATCGATC
 CAAGGCGACCGGAGCAAGCAAGTGGTTACAGAACTACGACCAAGCAAAATTTCCAGGAGTGGCGTTCCAGGAGTGAAGCAAACTGAGGCGGCTCAAGAGTGGAA
 AGGCGCAGATCGGATAGAAAGAGCTCGTCAAACTTGACTTGACTTCTGGCTTAAAGAGCAGCAGATTGTCAGCGAAACCGAACGCTATTTATAGCCAGCAGGGGCGATC
 GTGTATCTTGAAGAGACCGCGGAATAATCAATATTGAGTTTATATTAATCCACTTAAATAATAGAAATTAACAAAAATTTGCCAATTTATCAGTAATTTTATATCTCG
 CAGGCTATTTATCGCCATCTTCTGTTATACGAATACGACAACTTAGTATTAAGCAATATTTCTTACGTAATTAAGCAAACTGAGGCGGCTCAAGAGTACCAG
 CCTGAACTGAAAGTCATATTGATGAGGACCCACATGCTTGTCTACACAGAGGCGCTCAAGTCAACAAAGACGCGTGAGAGGAAAGCGGATGAGATGCGGATT
 CGGATAGGAATGGCGATAGGATGGGATGTCGGAGCCGAGGATAACAACTACCGTGCAGACGTTGCGAAAGAGTCTCCGCGGAGGCGAGCTCGAGTGGTCTCAAGCC
 CAAAGGATGGGCTGGGCTACACAAAGATGCAAGGCTGCGGATGTTGCTCAGCTTCGGGCTGGGAAAGTTTAACTGCCCTGAGGAGGAGGAGGATGCTGATGGT
 TATAAGCTGAGGCGCGCAATCCTTTGAGAGAAAGTGCAGAGTAACCGTAGAATGCGAAACATATATACGAACCTCACTAACGATGGGTGAGAAATACAGCTGGAAGCCA

FIGURE SHEET 8

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AGCAATTACTTCACTTCAAACCGAACAAGTCACATGTGTAATGGGTTCAATGGGTTGAGCACCGAAGACAGTTAACTCATCATGGAATAAAGAAACAGCCCGCACAA
ATTCTCTTTTATATGCGCACTCAATAATTTATATAGCTGTTTAAAGAAAATAATATTATCAACAACTCAGGGCCCCATAAAAGAGTAAGCCAATAATTTATGTACTTATCGCC
TTACTTCAGCTTCGCAAAAGTCAGATTGAACGCGCAGGAAGGCGATTGCCGACAGCGGATTCCACAGCTGGAGAGGGTGGCGCATGGTGTAGAAGGCCATGGAGCGATGGTCGC
ACCCGCAAGCTTGAACACCCGCTTGGCGATGGGTACCTGCCCGCTGTGCCACTTCACCGGCCGGCAGCGGGACACCGGAAGTCCAGGGCATCCGTGGCCCGACTAGTAGAGGG
CCACCGGTTCGACCAACCCGCGTGATTTGATGAAGAGAGTGGTGGCAACCGGATCTGGGCCGCTGAAGGAACCGGGCCCTACCAAAAAGAGAAATCACTGTGATGAAC
TAGATTTAAGAACCTTAGATACAAAGTTAAATGGGTGGATGATTTCAGTTGAAGAAGATAACTCACCAGCACTCCAGGAAGTCCAGGAGTGGCCATTGGCTGTGCAGAACTC
CAGGGCATTCGCTGCGGTGTAAATCGAATCCGTCACAGCTTCGACTGCTCCTCGTTCGCCGCCAGTGTCTCTCAGCGGCTTCAAGGCCAGGGCGATGGCCACACGGCG
TCGTAGTTTTCGGCGCGTACTGCGAGATGCCCTCCGGAAGAGAGTGGCTTCGCTGGTCCCTCCAGCGCCCTTCTCTCCGCCGACGAGTGCAGATTGGCTTCGACGCA
TACGTATCTTGAACCAATAAACCGAACCAATCATCTGCTGCCGTTGGAATGGCGGATGGCTTCGCCAGTGGGAGTTGAACATTGTGATTTCTCGGAAGTGGAAAGCGC
ATTAACCAAGCTCGAATGCCGCGAAGAAATGTCCGCCAGTAATGAGATTGGCTATGGCAGCTACCCTCTGCCCAATTTTCATGCAGCAGCAATAAAGAGCCATAAAACGCG
TCTGCGATTGGTAATGCGAATGCAGTACAGGCCCGCTTAGGGGGCTTCCGACTTGCCATCGCAATCACCATCGCCATCGCCATTGCTGAAACCATTCGCGAAACGCAATCGCAGC
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AAAAAATCGGGAATAAAAGTTTCGGCAATTCGGAATAATGTTGCTTGAAGCTGCCCTTGAGAGTTGGGTTCAATTCCAATTTGGATCCGGAATTTATATGCATGTATAACT
TAACAACAATCTATTCTTGAATATCTCTTTACGTTATCTCATATGCTGTCAATGAAATTTATTTATTTCTTCGCTTAAATTAAGTGAATGATTTTCTCGCATGCCACTCC
TCAACTTACAGCTCCACTATGCTAGCTTGTTCAGTTATTTTCCAACGATGCTGTGTGGCTTGAACACATGAGGTTTCTCCAGCGGAGTGCAGTTCGTGTTAGAGCAGCGGTG
CGCTGGTTCGGCCACCACGAGGCCCTATGCTCTCGTGGAGATCCAGGCGTAGTCCGCCCGCAACATTCGAAGCCTGTAGGCCCTCGCAGCAGGATTCGGGGGGCAGCTCTCT
GCGAGAAGCTGCGCATGATGATGCGCGTCTGCTGCTCTCTGTGGCGATACGAAACGAAACGAGCATGTGCCATTGTGCTATTGTGATCTCTCGATTAGCACAAGTGCCA
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GTCAGCAGTGAACCTCTCTGCTGCGAGAAAGTGGTCACTGCCGCCGACGCAAACTTCCGAATGAAAGCGATCGCGCGGGATTTGTGTAGGAGTCCCGGGCCCGCTCC
TGTAAGATGAGGGAACCTCCGCCGCTGCTGCTCAACGCGCGGATGTGAACCGGAAGGATACCTTCATCGAACGAAATCGTTAGACTACTTCCACCTGATCGGGGCGCTT
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AATGCTCTAAGTGAATTTGTGGACCTCATAAATTTGTGGGCTGTGGAATATTTATGGCTCAAAAGTTTATGGGGTGTGGAATCGGCTGGAATTCGGCTGGATTTATGCTA
CGAGCTAGCAGCTGAGTGTATTAGATTATTTGAGAAATTTGCCGATACAGGAGTGTGGTGTGAACCCGAGCGAGCTGGAATTTTGCATTTTAAATCAGTCCGAGACT
CACCTGACAGATGTTCCAGTAGGGCACCACCTTCGCCAGGCTCGGCTGACCTCCGAGCAGGCGCATCCGACGACATCACCATCCTCGTCGAGGGCTGTGTGTAGTGGCG
TGGAAAGAGCATCCACGCTGCTCCAGTCCAGGATCACACTGATTTCTCGGAGAGAAAGTTGTGAAGAAACGGAGAGCAAAAGCACACCATCAGAGATGAGTAATAGAGGGCG
AGGGAATAAGCAGTGGCAGAGCGGCTGTGCTGCTGACTCTTTTCGACGCAATAAAGGCAACATGATGGCGCAATAAACATTTTAAATTTGTCGACTTTTAGAGAGGCAC
AGCCGGAGATCGCGCTCCAGATATGACAGTCTGTTTAAAGTCTGATTTGCAATGATCATGATCACTTAAAAAATCGTTTGAACCTAGTTTCAATGAATGTGTGACATCAGTCTG
AGGTATTACGTTATGACGGTTTTAATTTATCGAAGAACTAACTATGATGCTTTCTTTTTTTTACCACTAATTAATGTAGCTTTTAATACACTACCTACCAAGTTTAA
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AGTTTAAATGTTGACCGCGCTACTGAAATGAATAATCAATTTTCGTAATGGAACAGCGACGCTAGAAGTGGCCCAACGTGGGCGATTCGCTTGTGCAACACTGG
TTCTCATCTGAGTATGCTTGTGTCAGCAGCTCGAGGGTGTAGCCCGGACAGGCGCTTGGCGTTGATGTGCTCCAGCGCATTTGGTGGCAGCTCCCAATTCGCTCAGACCATC
CGGACGTGGTCCCCGCGATGTGGACAGCTCAAAGAGTCCGAGCAGCAGATTTGCGCATGATTGGCGTGCATTTCCGCCCGCGACATCGGGCGATTCGTTTGTCAACACTGG
TGACCTTCAAGTATGCTGCTGCTGGGGGCGAGTTGAGGAGGTTGGGGCTGGAGGCAATTTGCTTCCGGCAGATTGCTGTGCAATTTGAGGCGCTGCTGCCGGCTGT
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AATTGAGCGACACCTGGCCCTAATTTGCCCTGTAGCCAGGGGCTGCAAGCTCGGCTGATGACTTCATGGCAGCGCTGCCAGCTGGCCAGTGGCCAGGCGCAGGCTAGGCTC
AGTCCGCGCGCCGCTATGGACCTGATCTGCTGCTTGGACCGGTTGAGGACCTGTCGCGCATTTATCGGCATCTTCCGACAGGATACGCTTATCGTTCTCTTCC
CTATCAGAGGCTGACGAAACAAACAAACAGCAGCACAGCAAGCAAGCCGGAATATATCGCTTTTATGACTCTGCCGACAGGATACGCTTATCGTTCTCTTCC
GACGAGCGGCTCTCTCTTCTGCTCCTCTCTTGTATCGGCGCAATTTTCTGATCGCTTTTATTTGGCCTCACTTTTGTCCGATTTTGTATTTTGCACATCACTCTGC
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TTTTCTGATAGAAAAAGCTTATTTTATATGCACTGCTTTATAGGATTAACAATTTAGGTGAATTTTATAGTATTTTGTGTAGTATGTGATTTTAAATGT
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CACTTGGCCATTTATTTTATGCTAATGGCAAACTTTTGCCTTTTCATTGCGGGCTTTTGGACTCATTTTTGTCCAGTCTACAGCTCTGTTATGTGCTGCTG
TTTCTCTGATGCGCGGTATGCAATTTGTTTCAAGATTACCACTTGCAGTCCCTTTGCCGCTTTCGCCGCTTTTCACCGCCCACTTGCCTGCACCTAACGCCCACTTCAT
AGACGCTTTCTTCGCGCTTCTGCTTTTCCACAGCAGCGCTTTTCGCTGT
(SEQ ID NO: 13)

Exon: 8795..8184
Exon: 7427..7284
Exon: 6893..6618
Exon: 6085..5834
Exon: 5246..4884
Exon: 4786..4485
Exon: 4114..3976
Exon: 3067..2792
Exon: 2683..2365
Exon: 2235..1001
Start ATG: 8795 (Reverse strand: CAT)

Transcript No. : CT9836
ATGCGCATAATCTCAACCGGTTCAACAGGGACAGATCGGTCCATGGCCGGCCGTGGGACTGAGGCTAGTCTTGCCCTTGCCCTGGGCAACGTGCGCAGCGGCTGCCATGAGT
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TGCAGGGGGGACTCCGATTCCACCGCGCTCCGATTGGAAGTACAACCGACGAAAGTCAAACGCCGGCAGCAGCGCCTCAATTCCGACAGCAATCTGCCCGGAAGACCAAT
GCTTCCACCGCCACCCACTCTCAATTGCCCCCGGACGCGATCTTGAAGTCAACACAGGTGTTCCGAAGCGAAGCGCCGATCTGCCCGGCCGATCGACGCCAAT
ACGTCAAAATCGTGTGCTCGGACTTTTGAGTGTGTCACATCGCGGGGACACGCTCGGAATGGTCTGAGCGATGGGAGCTGCCCATCTGGCCGTGAGCACATCAACCG
CAAGCGCCTGCTGCCGGGCTACACCCTCGACTCTGTGACCAAGCATCATAGTGTGATCTGGAGTGGGCGTGGATCGCTCTTCACAGCCCTATCTACACAGCCGCTCGACG
AGGATGGTGAATGTGCTGGGATCGGCCTGCTCGGAGGTACCCGAGAGCCTGGCGAAGGTTGGTGCCTTACTGGAACATCTGTCAGTGTCTTCTGGTTCACATCGCCGGCGT
TGAGCGACAGCGGGGAGTTCCCTACTTCTACAGGACGGTGGCCCGGACTCTTCAACAATTCGGCGCGCATCGCTTCAATCGGAAGTTTGGCTGGGGCACGTTGACCAC
TTTCTCGCAGAACGAGGAGTTCACTCGCTGGCGGTGAACAACTGTGTCACCGAATCGGAGCGGCGCAACATCTCATGTCGCCCGACCACATCACTTTTGGCGGCCACCGCAT
AAGGAGCAGCTGCTGCTACTTAGGGAGACGGACCGCATCATCATCGGACGTTCTCGCAGGAGTGGCCGCCCTCTGTGCGAGGCTACAGGCTACAGGCTCGAATGTTTCG
GGCGCGCATACGCTGGATCTCTCACAGAGCATGGGGGCTCCGTGGTGGCCGGACAGCGCACCGCCTGCTCTAACACGGAATCGAGCTGGCCGTCGAGAACCTCATCGT

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(SEO ID NO: 16)

Start ATG: 2500 (Reverse strand: CAT)

(SEQ ID NO: 17)

Start ATG: 239 (Reverse strand: CAT)

(SEO ID NO: 18)

12/89

TTTTTCGCCCAAAATTCGCTGGCTTTTAAAGTC TCACATAAAATGAAATTTTCAAAATAGCAAA TAACA TAGGAATAATCCTTCTAAATCGCAATA TAAAAACATATATACAAGGA
GTGGTAAATACAGACATTCATTAAATGGCAAAACGATTTATTCATACAGACGTCCGATCCGCATTAACCATGCTTTTTCACACTTTTATATACCGT TAGTATAAATAAAAAAAA
ACATTAAATCCCTAGTTTATTAATACCAAAATTTGTTGGCTGAATTCGATCTTGAAGAGAAGCTGACATTTTTTTATATATGCTTCCCTACTTTTCAGGACGTGGCAACATAC

[illegible]

14/89

Exon: 1001..1278
Exon: 1327..1601
Exon: 1661..2187

15/89

Transcript no.: TC18507
ATGTTACTTCTCGCATCGATCTTAATATATTCTTGTGCTGAATCTGCAATCTTCGAGTGCTGAGATTGCCGACTGCAGTTTCTATGATACCGTTGATATCTCGGAAGGCCAGA
GGCTCTCGAATGGATACATACCTCTCGAGGAGTACTTATCCGCCCATTTGACGCCGCAAGTATGAATTCAAGCTCTCGGCAGATGGAGATAAGGACAGGAGTCCCAAGTCA
CGTCGAGGAGTATGTGTGTCAGCAAGCAAGGACTCGGCTCAAGTTTGTCTGCCCAATGACCATATGATATGGATATGGGCGCACTACGCCAACATAGACACGAGGAGAACCAAG

TTACTGGACCCCATGCTAAATGTGACCCGTGGACGACGGATCAGTGGTGCAAAGGCCACTATAGGAAGGAGCTCATGGTGCAGTGGGACTGCGCCAAAGCCCTGCGATGATATGTTTACCTAGACACACCGGGATATATGGATGAGTACACCTTTGTCGAGAATGGAAGATTGCTGCGCCACTACGATCAAGTGATTGGACAGTCGGGATACCTGCTCGCAGCACCGGAACATTCCGGGAGTAAACACACCTCCATTCGATTATACCCACACATGCTTGATACATGCCATCCGAGAAGCGGACAGACCGTTGTGATGATACATCGCTTGATATGCTTGCTTACCATCGCGCTGACTCTGCTGCTTGAAGAAGCTAATGAATTTGGAGGGCAAGTTGCTTATCTGTTACATGATGTGCGCTATTTTTCGGATACCGCTTCCCTACTCCTTGATCTGTGGGATTTATCGCTCGATTCTGTAAAGCAGCGTTTCTGGGATACITTTTTCGTCATGGCTGCGTCTTTTGGCTACTTATCATCAGTCGCGCACTATGGAAATGTCCTTACCAATCCCTGTGCCTCGATGAACATTTCGTCAGAACGTGCGTTTCTGCTATACAGCTGCTTCGCCCTGGGCAATGCCCTTTGGCCTTGACCGGAGTCACCTATCTGCTGATACATGCTTGTGAACAAGGAGGATGGCAGCCTCGCGCTGGGCGACGAAGGCCATGCTGTGGATCTATACATAAGACCTGGTTCGCAATGGTGATCTTATAGGACAGTATGGTCTCTGATTCCTCAACATACATATGTTTGTCTGACGGCTAAACACATATAGATATCCAAGAGACCTCTGCAAGATTTGCTGCCAATGAAGGTAGAATACAGAAGCTGAATTCGGACAAACAGAAATTACACACAATTTCTGCTGCTCTTCACTGTGATGGGATGTCGTGGAGCTTTGAGATATTCTCCTACTGGTGCACCGCGAAACCTGTGGGTC AATATCTTTCTGGTAGCTGACTACTTCAACTGGTCCCAAGGCGCTATTATATTTGTGCTATTATCTTGCGGCGCAAACTCTTGTACTATTCAAGAAACAGTAA (SEQ ID NO: 29)

Name: mth-like 2
Classification: G_protein_linked_receptor

Transcript No. : CT17758
ATGTTGAGCCCATTCGATTGGCGCGGGGCATCAGCAGCAGCGGAACCGGCGGCACCATGGCAGCCAGCCACTGAGCAGCAGCGCGGCAACAACTGCGGCGGCAACGGGAG
CAACTGCGGCAACTGCAGCAACGGCAGCAACACATCGGCGACCTTGAGCAGACCGCTGCTTCCACCTCGACACGGCAGCGCCATCCGCGGGAGGCACGTGGATAAATCA

TTGTCAGACATTACTGAATTAATAATAACTTTAAACATGCCACCTAACAAAAAGCGAAACAATTGTTTAATTACATTGTGCAACCTGTGGCATTAGTAAAAAGCAATATCT
AAGCAAAATAAGTCGGTGAATTTTGATATGCAATATAGCCAAACAAAAAGTCGGTGTATTTAAATATCATGATTACGCCATACGTTATAGATGTTTCATGCTCGCTTCGTTGT
TGATTTCCGCTGATAGAAGGACATACACTTTTAAATCTTGCTATGCTTGGTATTTTAAACGAGCATTCGAAGGTATCTCTGCCCATTTCAAAAAGAGCTGAAGCTCTTTGGC
AGCCAAAAAAGCTGACAAATTTGTGCCGGAAGGAAACAACCTTCCATTTCATGATGATTATTGCCAATGGAATAACAGGTTGGCTTCAGTTGCGTATGTCCCTGCTATTTCGA
TTTTCAACGAAATTTCCGTTGGCATTCACTCGGAATGAAAGTAAAAAGAGTGTCTACTTACCTTTTGGAAGCCCCGGAGAAAGAACAACTTTCAATTTGGCCAA
TTAGTGTATTTTAAATGSCAAGTTACCGGCTTTCAAAGGCCACATTTTGTTTTGAAGTCCGGAGTTGTAATGTAAATGAACAAATTCGAGTTTCCATAGGCCA
ACCTTTACACTGAGTTCCTTTCCAGTTGCTCAAGTTTGTCTCTTGGCAGCTGCACCTTGTGTTCCTGTTTGGCAGAAAGTTAGTTTCTGCCAAGAATTTGTCTTTGGCCA
AGTACTGCTCAGCTCGGTTCTTTTGGGCAAAAGTTGGTATTTAGATTTCGACGTGGAAGAACACTTCGGCGAGATCTTCAAAATCTGCCAAACCCGCGCGGCTCAGCTGGC
CGCTTCTAGTTATTTAATAAGTAAATTAAGTTCCCTAATTAATACATTCTCGCAGAGATTACATAAAGTATCTACCGGCAAGCTTCTCTCTCATTTGGCGGCTTAAT
AAATTTGCAGACCGATTGCACTTTCCGAAACTTGATTTGGTTTGGCTCTGGCTGGCAGTCTGTTAAACAAATCATGATTGCTGAAATGGTCAACTTTGGGCACTGCACATTCGTA
ACACTTGAGATTAATAAATTTCTTAATAGGGGAATAATACACTTATTTATAGAGTATTTTAAATATATACAAATTAATCTAGTCAGCGGCTTTGACGTAATTTATATTTAG
TTCGAACAACCTAATTTGACAACATCAAGAATCCAATATTACGAATTTTAAACATATCTACGATCGGAAGGGGAATCGAATTTCCATAATTTTAAAAAGGAATCGGAATTT
TACATTTTCTCGCGCTTTTGGCGAAGAGCGGTGATATGGTGACGTATTTATGAAATTTGCTTTTACGCCACTTCCATGAACTGAAATTTGGTTTCTACTGCTGTTTGTGG
TCGATATATAATGCTGAGATATATATATGATATGTAATGCTTTCTATATTTGTAGACATTTACGAAGGGTTATAGTGTGATATAAAGAAAGAAAGAACTTGAACAAAC
TCAAGTATATAAAAAAGGTTTCAATAAAGATATAAAGTCTATTAGCTTACGTTACAGGAGCATTAAGTTTCTTGTAAAGACAGCTTTATCGAACTTTATCTTATAGAT
ACATATATCTATTACTAGGATAAGTAAGTAATTAATTTATGACAGCTGCATGACATATGCCAAACACTTTAAGATATCCCAACCGGCACTTGCATACATTTATC
AGCAATTTGAATTAATTTTCAGGTTCGGGTTACGTTAGTTTGGTTTGTAAATACATTTAATGTTTAAATATCAGCAGAGAGCAGGAAATAATGTATTAAGTTTCGGGATTG
TGTTAAACTTTAAGCGAATAGTTTAAACAGCTGTTATTTGGTTACCCGCTGCTTTTAAATAAATTTTCCCTTTCGATGCCACACCACTGAGTTATTTCTTAAATGGTCTGAAA
AATAAGAGAAAAAAGGTTTGGCTCAAGTTTATTTGGTTTCGGCTGCTTTCATCAATTTGCTGCACGCAAAATGCTTCTCAAAAAAAGAAAAAAGAACTACTCTTCTATTTA
TGGCATTCTTTAGTTTGGCAATAAATTCGGATTTAAGCTGGTTGAAAAAAGCCCATCCTTTCAACGGGTGACTTAAACATATTTTAAAGCAGACCGCTCAAACTCAAACTGGGT
GGTTAATTAATAAGACCATGTAATAGTCAACTGCTTTTATCTGCTCCACTTTTAGGCTACCCCGTGGACAGCTAATTTGCAAGTACACCCCATTTAGCAAACTTGAACAAAG
GAAATGATCTGTGACCGGTGCACCAACTTGCACACTACCCAGCAGCATACGGCCACCACTTCCACTCCAGCAGCAGAAAGGAATCCGACCGCTGAAACTGAAT
CTGAAGGCACTCCAGTGGCGGCGACACACGGAATCCTTGGCCAATGCGGCCAATGCGAGGCGCTGGCAAGTAAATAGCGTAATCCTTTGCGGGCAGCTACACCACTACT
ACCTCTGTTCCGAGTAGCTCGTAATATTTGGCCAACTCAGTCACTGGAGGCCATCATCATGACGACCTGCCAACTGCGAACGGATGAGGCTGAGCAGCTCTTGGCT
AACCGGTGCCCTCTCGCTCTCCGAGATGTTGGCCAACTGAGCCACAGCCATTCCTATGGGAGCAACCTTCAGCGCTGGAGTTTGGCCACCGGAATCTCCGCTGTGAAT
TAGGGCAAGGACACGACAGCAAGCTGAATGACAGCGCTTCCACGGGTCTGAGGTAAAGTGATGTGCTTAGGCTTTTCCATTTCCATGGCAACACACACAGATTTCCATTA
AGCTCATGACTCGGGCTCAACAGCTCATCGCGATAAGCTCAGCTTCTATTTCACGAGATGCATCTCAACTAATGTGCTTAGAATCTACTATTGTATACATAACATATA
GGGAATATAATCAGATTAATTAGAACTCAGAACAGTCAATACATCACTTATATAAATGAATAAAGATCTATCTGAAACTGAAAGAAAGAGGATCATTAGCTTGGGT
ACAACATTTATAAGAAATTTGGCTTATGAAGTGCRAAAAGAAATCTTAAAAAATCAATCAATGTCAATAGTTCTGAACCAATAAATTTAACAATTAATAATTTAAAT
TATTTATATGATTTAATTCACACTTTTAAACATTTTACACTACTCGAAGGAGAAAGTATCAGATGACTTGAGCTATACTTTTCAAACTTACACTTACAGCAATCTAA
CCAAATGAAAACTTATGAAACCCATTTTAAATGAATAAATGAACAACTTACTTTTCCATTTCCCGACATACAGATTTATCCCCAGCTAGCTTAGACAGGGCA
AGTACGACCTGAGCTACATTTGCCAAGGTGAATCCCTTTTGGCTACAGTTTCGAGCCACTAAGTCGAGACCTTCTCTGATATGGCCGCGCTGTATGGCTGATTTTCGGTGT
GGGATGTGTGGGCAATCGTTTCGTCATTTTATGTTGGCCAGGCGCAATCTCTGGGCACTCTCTGGCAATATCTCTGGTGATGAATCTGGCCACTCTGCGACTTCTGATGCTC
ATCAAAATCCCGATTTGCCATTACAATAACATCAAAAGGCGCAGCTTTTAGGAGACATTTGTAAGCTTAAAACTTGGGGAATAACTTACATAATCTATATAATTTGAAA
TTCATCTCGGCTTGCAATTTATGTCAAAATGATCAACAAAAGCAATCTACATATAATAAATAAATTTAAATTTGGCCAAAATGCTTGGAGCTGATTTTCAGCTGTTTCTGCA
ATTCAATTTGTGGGATCTCTTTTCCGAAAGCAATTAATAGAATTTAAATGAACAAATAAAAGTTCAATTTGTAGCTTACACATCTTGGTTGATGAAACATATTTCCAAAA
TGAATGGCTATAAATTTTCCACGAGCCTTTTGTGTGCTTGAACGGGTGGTTCATAAAAAATCTACTGACTTTTCCGTCCACAGCCTTGCCTCTATGATTTGTGG
GTGGCCTAAGTGGCACTGTGTCATCGGTACCTTACCAGCCTCGCTTTGGATCGGTAGCAATGTGGTGGTGCATCTCATCAACCGCTGAGACGCTGTCCCGCTCGCAT
CTACCTGATCATCTCTGCTGATCTGGTGTACAGCTTCCCTGTTCGCTGTGATCGCGGCTTGGATATCGGACTTCTGCTGTGCGCAGAGGCTTCTCTACCACCTGCAGC
TTCGATTTCTGAACAGGAGATGCGAGCTGCATTTTCATGCGACTGTTCTTGGCTGCGCTACTGCACTTCACTGACCTCCATTTGTGTACTCTCTATATATAGTA
AGTGTGTTCTCACAGCGAGTCGAATACAATCGAACAGGATAAGGCCAAGACGAGCAGAAATTTGGCTTATTTGGGCGGCTATGATTTGGTTTGTGGTCTCTGGCTGGT
ACCGTAGCCGCACTGTTGCCATGATGGGTGTTCTCGGCTGGAGGGACATAACCGCCACTGGGATCGATGATACCCGCGCTCTTCTGCAAGACAGCTGCTTGGCTGGATCCC
TACTTGTATCGGCCGCCACCTACCGCGGTTCTGCTGAGGTGCGAAATGCTCTTACAGCAGCGGAGGTGCTGCAAGGGCTTCCACAGCAGATCTGCTATGATACCGCAT
CACGATCGTCTGTACCCACCGATTGAGGACGAGCACCACCGGTGAGGGCGGAATGGTGATCACCGCATGGAAACATCTGATGAACACATCTGATGATGGTGGCCGA
GGAGCGGAGGAGAACGAGGAGATCGTTGTGTGGCGGAGATCAACAAATTCGATTTAGCAGCGTCAATGGAACAGAGTAAGTTCTGAAGCTGACGATTTGCCGTTTAAGAGA
ACGGAATCTCTGTCAACTGTGTTGCAACCACTAACATAACATTTTAAATCGATTTTAAAGTTTAAAGTTTACAGAGCTGTTTATATAATTTGAAATAGTCAAACTTATGATCAC
GATGATCTGAAAACCTTTGCTTGAATATCTGAACTTCGCTGTTTAAATTTTCCATTTTCCAAATGAAATTTCAATTTGATTTTTCGCCGAGTAATTTATAAAAAACCT
TCCATTTCTCGAAAAAGTAACTGTTTCTGCTGTAAATGAAATGTTTGTGCACTGCATCTGAAGGCATCAAAATGGAAGCTATCAATAATTTGAATGGATCGGAC
TCAAGTTTCTATTGTGAATACTAATCTAATTTGTATGTAGAAATTAATAGACCGAACTGAAATCAGCAATGGCAGCAAGTAATTTTCGTCCGCAATGATATGCGCAAT
GGCATTTGTTCTATCTGAGTGAATGCGTTGTCTATGATGGACCTACAAATGAAATGAAATATAATAATAATAAGAAATGAAGCAGGGAATTCGCCGAGCAGTGGGAAA
TTGGCAAAAGGCACACAGCAGACAGTGGTGATAGTAATCAATTTAATGAGCTTTATGAGCTGTTTAGACAGGGGGAATGGGATCAGAGCAGGGAATGGATTTGTAGCC
TATATACATATATATATACACAGCATGTGATCATGTATAGAACATCGGAATATAGAGCGCAATTAGAGAGGCAAGGCGTAAATTAGTACAAAGAAAGAGGTTGGCTGGG
TAGGGGCCGAAGTGGAAATTCAAATAGTCAGATATACATCAATTTACGGGAAGTCACTCGGACCAAGGTCAGCCGTGGTATTGGATGTTAACAGCGGATGTGCAGATTG
TTGCAATTTGATGTTAATATGGAATAAATCACAATCAAAGCAGCCTCGGACATCTTGATTTCTGAAATGGAGTC
(SEQ ID NO: 34)

Exon: 1001..1464
Exon: 7788..8022
Exon: 8083..8341
Exon: 8920..9245
Exon: 9609..10501
Start ATG: 7845

Transcript No. : CT17820

CTGGGATGCAACGGATGCGGATGCGCTTAAATCCCCGAGCGGTTTGGCGGTGTTATTGAAAATTGGGTATTCGCCGCGTATTACCAAGTAATAGCTTGTCCAAAGGCCACTACG
CTGTCAAAGGACGGGCTTAAAAACGCTACTGCATTTTTCAGCTACTATACATAGATAAAATTTTCGCGGTAAATCGCGCGAAATTGAGTTAAAGTTTGTTGTGTCGGGTTGGCGGCCA
TTGGGCAATTGGTGCTGGCCAACTTTTGGACGGCAAGTAGACAAATGAAAGTAACCTTTGTGAAGCTCAGCAGCTTACCAACAAATTCAGTGTGTTGTCGCGAAAAAAAT
TAAAAAATTTTGAATTGAAGAGGACACAGTTCGGTGAAAAAAAATATTTGCAATTGCACAAATGAAAAATCAGCAAAAGTGACCTCCAAGTTTACGAAAAACGAAATTAACCTA
ATAACACTCCCTCGCAGGGTCACCCGGTGGACAGCAATTTGCAGTAAACCCCAATTTGGCATTTATGACCAAGCGAAATTCGATCTGACCGCTGCAACACCATTTGCACCACTACCA
CCAGCAGCATCAGCGCCACCCTTCCACTCTCAGCAGCAGCAAGGAAATCCGCACGCTGAAACTGAATCTGAAGGCATCCGAGTGGCGGCGACACAGGAATCTTGCCTCCAA
ATGCGGCCCAATTCGAGGGCGCTGGCAACTCGTAAATATTTGGCCAAGTCAGTCATGAGGCGCATCATATGACGACCTTGCCCAACCTGACAAACGGATGCAGGTTGACAGCAG
CTTCTGGCTAACCCGTCGCCCTCTCGCTCTCCGAGATGTGGGCAACTTCGACCAACAGCCATTCCTCACTGGGACCAACCTCTCAGCGCTGGGAGTTGCGGCGCCGAATCACTC
GCTGTGAATGTAGGCAGGCCACACAGCAAGCAGTGTAATGACAGCGTTTCGACGGGCTGAGCAATTACAGTAATATACCCACCTGACTACATCTACTACGGGACCAAGTAGCCAC
TGAGCTACATTTGCCAAGGTGAATCCCTTTTGGCTACAGTTCGAGCCACCTAAGTCGAGCACCTTCTGATTATGGCCGCCCTGTATTGCCTGATTTCGGTGGTGGGATGTGT
GGGCAATTCGGTTCTGCTATCTTTATGTTTGGCAACCGCAAAATCTCTCGGAATCTGCGCAATATCTCGGTGATGAATCTGGCCATCTGGCCACTTCTGATGCTCATCAAAATGC
CCGATTTGCCATTTTACCAATAACATAAGGAGGTCACGCTTTAGGACAGATTCGCTCTGCGCTCTATGATGTTTGGGTGGCTCAATGGTGCACCTTGTGCCATCGGTATCCCTC

19/89

Start ATG: 522

Name: Rhodopsin 3 and 4-like (Rhodopsin 7)
Classification: G_{protein}_linked_receptor

[illegible]

20/89

TTATCTTTGTATGTACGATTAAATTTAATCTCTAAAGAGTCAAAGTACTTCAATAATTTACAAATCTGTGCGAGGGAATGTAAGCTTGCCC AAATCAATGGAAGCTACAA
TCGTTGTCAGATGGAGATATTTATGTTAGAGGAATATCTCGGATCGAAATATTTAAATTTAAATTCCTTTTCATATATTCTTTTAAAAAATAATAGTCCTGTATGT
ACAAATTCGAATACAAAGGTAGAGTATGTCCTTAAAAAGAAAATATCTTTCCGAATATATGTATATATATTAATAAATAGGTGATACATACATTTTTCGTAAGTATCCT
AGAAGATTTGCCACAGTATCAGGTTTACAAAAAATGATTACCTATGTGCGAGTAACTTAAACGGCAGTGCACTAACTAATCTGATTTGTGAAATAATGCAGATATAAAACATT
TTTAAATAGAATATATTTATACATCTGATTTCTCAACATCTAAGCACTCGTATAGAGTGCTTTGAATATATTCTAAATACATTGGCATGATCCAAGTTTAAATCAGCCAA
AAACTGCTCCAAAAACGGAAGATTCTACACAAATCGCCTCGCCCTTAAAGTTTCTGCCAAGTATTGGGTGCGTGCGGATCAGTTT
(SEQ ID NO: 37)

Exon: 2..134
Exon: 208..388
Exon: 2321..3216
Exon: 3631..3664
Exon: 4521..5472
Start ATG: 2

Transcript No. : CT18539

[illegible]

Start ATG: 1

MRNSTNIFLTNLISIADLLVLLVCTPTVLVEVNRTPETVWLGHMCKAVPFVELTVAHASVLTILAI SFERYYA ICEPLKAGYVCTKGRAILICVLAWGIAAFTSPILWVAE
YKLEAYIDGSSVAVCLTQA ISDWTLA FLMLT SVFFVVFVPTVLVLYG I IARNLVSNRAAMLRAPRTKPELSLKARKQVVMILGAVVLSFVFCLLFPFVUTLWIILSTDQTL
HDLGLVRRYSLLYFCRIMLYLNSAMNPILYNLMSTKFRRGFKRLQDAGRLLLELVLTLGRKKEDSSRGRRTLSLGMGTNTNTNTNSNATGATSSSILRSSNNRCSDEIS
RTRLKIENMQMPCGSDLEAMMLQHS TLGKGIARRVSDSLMPLRLNRHQPRRHKQPSIFDESELEENKRLGVQTSKQKQPIAALSTYSVEPYGDDGTAPVCTAADGWSFI
YFGVCTITVFFFLPFGILVLLYAAIYKLLRPNNASHRPTSPQCPQSGGATSGSSQVPTKNSHQSGNRMKRRKGIVIFMLVAVVSFVFCLLFPFRAFTLWLIILASAEDV
GLGIAGYNYLIFYSRFMLYLNSAMNPILYNLMSSKFRSGFWRLLLTCLGQRPHHHHRHHYHQRHQPTAGGSGRNASTRQEQDAEEGAALAGTTSARHPRTLRREATFLINS
ITSSGSDRTTSSSAWRNSLSISLSESERGILGAAI GTTAAATVTTACLQERRASKT *
(SEQ ID NO: 39)

Name: Thyrotropin-releasing hormone receptor-like
Classification: G protein linked receptor

Calera Sequence No. : 142000012807287

[illegible]

21/89

CACATGACGATCGCCTCGGAGGTCTATCCGAGCTTCCCTTCAGGTGTTCCGTGCCAGCGCAATCTGCTGGAGTTGATTAACACTACTCGTGACCTTCTACATCTACTGCC
 TGTTACGCGAGGATTTCCGCAACACCTTGTTAGGACGATCAAGTGGCCCTGGTTGAAGGGCAAGTTCTGCCACCGAGCGGAGCAGAGGTGAGTGCCGATCCACCGGCCAC
 AGCCGGAACCGTAGCGGTAGCCGGAACCGGAAATGGCCAGCTTTCATTTTTCATCCGCAATACCGCGGTGACTCTCACACCAGCAACCGCAGAACCGCCCGGGTGT
 GCGCATGGCGTCTTCACTAAGGTGGACAGGGGCCATCAAAAGCACCACATAAATACCACCTCGGGCCTGGGACGATCGAGTAGCATTAGACAATGCCCTACCACCAACAA
 CAACACCACCAACAACCATCCGCTACGATCGGCAATCCGAACACGAGAAACCTCCCAGCCAGAGTAACCCACCACCCACCCCAACACTCCAAAAACCGCTCATCACCAC
 TCTCCACTTACCCCCAGGTCGAGTCTAAAGGGTGCCCTTCTCTCGTTCGAGATCTCGCAGCATTTGAACGTGTTCAAGTAGATCGACGAAGAAATGATATACGATA
 AACAACTAACGATTAAAGATAATTGATATTACGGCCGACGATTCGGTATATAAAGTGCTCTGTGTGTTCCCTAAACATATATATTAAATATATGGATAAACACTTTTTTT
 TTTTATAGCACTAGAATTAGGTACTTAAAAACCAAACTTAATCGAAAACACATCCGTAATTGGTGAATAATTGATATCGATAGGTAACCAAGTTCGACTTTCGATCAGCAT
 TTGCGCCGAAATGATTTTCAACCAATAGTCTTCGGAATGTGTATGTCGAGAGTCAAGATCCTCAAGCCACACGATAGATGCAGATATGATAGTTTATCAATGGATAGA
 TTGCTTGGCGTATTGATAACCAAGAGCTGTTTACTCTAGCATTAGTGCAGACAGAAACAGGAACAGCGACAGGACAACGACAGACAGACATATTGACATATAGACAGT
 ATAGACAGTGTAGATGGAAACAAATAGCGAGAGAGAGAGAAAGAGTTTACGATGATAATATGCAGACATACAATACTTATATATACACTTGCATATGCCATATATGTGAATG
 ATATTTATGTAAACGACGTATTTATGTATACACGCGTAACCTTAAAGCAATATATTTATATATGCATATATGTATCTATTGATTATAGCATTTAAGTGCAGCGCAAGAGGAAT
 CCGGATTCCAACC
 (SEQ ID NO: 40)

Exon: 1001..1088
Exon: 1280..1429
Exon: 1851..1944
Exon: 1996..2821
Start ATG: 1001

Transcript No. : CT18637

TRANSCRIPT NO: 13509
 ATGGAGGCGAGACGAGGAGGATGAAGAAAAAGGGAACCGGAAAGGAAAGCCATTCCATTGGCCCCGCGACTCAAGTGCCGTTGTCTACCTTACTCCACGCCGCCCTGCTGG
 CCAATCGTGTTCGCCATTCCCTTCGGTATCCGATGCTGCTCCACAGGATCGAGGACAGTGGGAGAGTTCGCCGCCAGCCTTCTACACGCCGCCACCTGGAGCTTTATCTGGG
 CAACGGCTGTTTGGGGCTGGCGTAATGATGTACTAGTCTGACCAATTGAACGTTACGTGTCCGTTTGGCCATCCAGGATTGTCTCGACCAATGATGGGACCACTGGCGTT
 TGGGTATTCCTCACCTGCTTGGCCACGGTAATCGTCTATCTACCGAGCATCTTTCGTGGCGAACTGATCAAAATGCATCCTTGGATCGAGTGACGATATATGTATATTTCGAC
 CGCAGACCACTATCTACCGCGACAGCACTTCTCATCGCTCTACAGAATGATGATCGTGTGGAGGAGTACTCTTAAATCGTGGTCCCACTTTGGTGATCGGTGGCCCTCAACATCGGCAT
 CATGATGGCTCTACGGCGGACCTTCGCGAGCCGCGCGCAGATGTGGTGCATCGTCCGACATGCCACGGGACATGGTCAACGACATGGGACGATGGTCTATGGACATGGA
 CATGCACATGGACACGGCTATTTTGAAGGACGACGATCCCGGAAAGTTCGCCGAGGAGCGCGCTTGTTCCTGCTGCTCGCGACACATCGAATTTGTTCCTGGTCTCGCTGAC
 CGCGGATGGGCCATTCTCCACATGACATCGCCTTCGGAGGTCTATCCGAGTCTCCCTCTCCAGTGTTCGGTGAGCGGCCAAATCTGCTGGAGTTGATTACTACTACGCTGAC
 CTCTACATCTACTGCTGTTTCAGCGAGGATTTCCGCAACACCTTGTTGAGGACATCAAGTGGCCCTGTTTGAAGGGCAAGTCTGCCACAGGCCAGCAGAGGTGAGT
 GCCAGTCCACCGGCCACAGCCGAACCGTAGCGGTAGCCGGAACCGGAAATGGCCAGTTTCTATTTTTCATCGGCAATACCGGCGCTGACTCTCACACCCGAGAACCCG
 ACGAACGGCCCCGGTGTGCCAAATGGCGTGCTTCACATA
 (SEQ ID NO: 41)

Start ATG: 1

MGRAAEVVKKREPERKAIHLAPRLKCRCLFPYSTAALLATVFAIPFGIRMLVHKDRQWEEFGPAFYTAHLEYLNGCLGVGMMLLVLTIERVSVCHPGFARPMVGPPGV
VVFLTCLATVIVYLPISIFRGELIKCILGSSDVVYLRDRNTIYQTIYFRVYKIMLEVIFKLVPTLVIGGLNMRTMMVYRTRCERRRKMVLRSRPAHQGHGHHGHGHGHGHG
HAHGHGYLKDDPRKFAEERRLLFLLGSTSILFLVCVSPMALLHMTIASEVYPSFPFQVFRASANLLELINYSLTFYIYCLFSEDFRNTLVRTIKWPWLKKGKFKCHQAEHVS
ASPPATAGTVAVAGTNGHVSIFHPAIPALTLTPAEPDERPRCANGVLH*
(SEQ ID NO: 42)

Name: G-protein coupled receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384543

AATAATCCACCTTCGAAGGTAACCACTCTTAATTATGCTTTAAAAATGCTGGATTATATAAACCCCTAAGGTAGTTCATCCCACCTTGTTTCAACTGAATCGACCCACAAATA
TGTTGGTCTGGTATCTATCTACCAAGTAGGAGCTGCTTAACCCAAATATGCGTTATCTGTAGAGATCCGCATCTCGGAGGAGAAGTCGAATCCGGTAGCTAGCTGCG
AGTCGGTCTTCTTCAGAAATGGAACCTGGTGGGTGGTGCATACCCGTCGGTGGCCATCAGGAGTCTCTGTTGTGAGACAGCGCGCTCGTGGTCGGCAAGGAGAT
GACCGAGTCTGCTTACCAGTTGATGGCGCTTTGCTCTGGTGGTGGTCCACAAACGAGATCACCAGGCTCTGTTTACGAATCTTCCGCGTTATCGAGACAGAGAGCTC
GGTGGTGGCGCCCTCTGGCCGGCCACATCCATCGAAACGCTATCCCTCTGCCAGCTGTCTGTCGAGACCGTCTCTCGCGACCTCGCTGCCCTCATGTAGGACCAATTT
AGTCTGCTGTTGACAGAGGAGACCTTGGAGCGATTGCTGTCTCTGGCAGAGCGGTCCGCGATCCAGAAAGTATCCTCAGGAGGCTCTCTGCTTGATGAGGAACTGACG
CGATCCGTGAGTAGTCTCTGCTTAAGCAGCATCTGCTCGCAGAGGTTGCTCTGCTGTAGCATATGTTGTGGCCGCCACCGGAGTTGCTCTGTAGATATTAGGACACCGCG
TGTGCCCTTGCTGCTCGGACCTCTTCTCTGAGCTTCTCTGTTGGTATGTCGATGACTTCCGGGTGCTGTGGTGGTGGAGCGCATGGSCAGTTGTCTGTAGTACTTAGTGGTT
AAGGTGGAAAAAGGTTTTTATCGAAGGGAACACAGATTTCTTACAAATCTTCATATCAATTTTGAAGGCCTTCACATATTTTATCAATATATTGAGGTTAGAAATTAACATA
ACCTGAGGCGCTCTCGCAGCAGCTCGCGAGCTTGATGATGGCCGGAATCCGCGTCGGTACTCTTCTTACATGAGTAGATAGGTTGGGTTGATGACGCTTCTCGAGGAGCCAG
CCACTGGGCCACCGGACATACCTTTCTTAGGATCTCGAATCTCTCTGGAGATGTCGAGCGCAATTTGATCCGCGAAGATGACATGAGCGCGGACCGGACGACAAACAA
AACAGGATGACCACGGCCACCAACATCTGTATGACCTTACCCTTGTCTTCTGCTGCATCGGTCACATTTGCGCGTCTTGGACTCCCGGGGATGGACGCTGTGGAGACCT
TGATCCAGAGAGCACTGACGAGCTGAGGAGTATCAGGACATGGGCAGGAGTAGCAGGACCCACGCTGGCTAGCAGGAAGTAGAGTTTCCATCCGTTGCCCGGGGGCCAGCT
CTCTCGCATGAGGACCTCGGGCTGAGGATGAGCCGAGACAGGCGCTCGGAGAGGCTCCAGGTGGGACCGAGGTCAGAGAGAGCAGCCAGGAATGGTGTCCACACG
GCGATTACCCATATGCCGATTATCATGATCGGGCAGTGCCTTTGTCATCTGCTTCAGGGGCCACCAAGATGGCAATGAOCTCGAAGAGGGGATCCCAACAAAGGGGATTT
TTTACCAAGTAGTCAAGAAGAGGAGTCTCGTAGGACGACGACGACGAGCAGGAGGCTCCGGTCTCGAGTGCACCTGACGGCTTTGAGCGCTCAAGGATATGGCAATGACCATCAA
GGTGCAGTGAGTCGAGTGCAGGCGACGATGACGATAGCAGACCGTTTCAACTAGGAGGAGCGACTCGGATCGCAAAATATCTCCGCTGATTAATTGCGACACGAGTATCCGA
TGCCAGCAGCCATCCTTAAGCCTTTTCTGCTGAGTGTGCTTTTCCAATCTCGACTTGAACCCGGCTGCTACTCTGCTCTGCTCTCGCCGCCAAAAGAGCGGGCGCT
CGGGATCTGCTCATTTTATTTCTGCTGCCATGAATTTGAATTTGAATAAGAGCTCGGAGGCTAGTGAGACGAGTCGATATGGGGCACAACGAGTTTTCAGCATTTTCTT
ATTTAGCTTAGTGCGAGCCGCTCATTTGCGCCCGCGCTCAGTGGCTATAGACAAGACTTGTCCGGAGCAACGAAACTGGTTGCTGCTCTGCGCCGGAACCATGTAAATTA
CGTGCAATTGTGTCGATTATTTCGACTTGACCTTGAGGTGTGCGCTGCTGCTGTCGCCGCAAAACATTTGCGAGATAACACAGCGGCTAATTCCTATGAGGGCGGCATGCCA
GGCAGCTACCCGAGAACCTTATCTCCACCGATAAAAATAGCCACTTAAACATGTGGCCGAGACCTCATCTCTCGCAAAAAAAAGAGTGGCGCGCTGGGCGACCAAA
CGGACCTGTCGCAAAAGCCAAATACAAATTTGCTAAAGCTTTATTTAGGAGCAAAACCAATTCGATTTGCATAATTTGCCATGCGACTTCTATGGTCAATAATCAGGCG
AGAGCTTTAATTAGTGTCTTTTGGGTAAGAACAGCGGAGTCGGAAGAAAGTATACGAGTTTTCGCTGGTGACAGATTAACAACTAACCTCTAATTTGATGTTGAATTT
TTGTTAAATCGTATTAACATACATTTTATGATGTAAACCAATTTTATGAGCTTTAAACAACTAAATTTTCTCTATAGTGTCTCTATACAGCTTTATGTATGATTTGTTTAC
AATAAACTCTCTTATAGATTCAATTTGTGTCGGAATAATTAATTTAATAATATTTTGGTAATCAGGGTAAATGTTGCCCATCAATTCGTTGAGTACCATATGCGTA
AAATCTTAAAGGACGTCAGTTTCAGTATTTAGGATATTTAATTAAGAAAGAACTTGCACAACTCAATTAACAAAGTAGTAAATCTCGGCTTTTAAAGCAATTAATAGC
CGCAAGGTGATTAATTAATTAATTTGCCACCAATCTGTGTTTCTGTAGATCCAGTGTTTATAGGCTCTTATTCGTTCTCTACTGCGGTTACCAATACGCACTGTTT

GCTTTTGTCTTGGCAGTCTATTTTAAAAGTTATCTTGACCATTGCAGAAAGCTATTAATTTTGATACTAAAGCTTGGCATTTCAGAACAACTTTAATGATGCTGCATCG
 TTATGTCTCTCTTGTGTATGTAGTAAAATCTCGGAAGAGGCCAGGAGTCTCGGTATATAAAATGCAGCTTACAAAATAACCGGAAACACAGATGTCTGTCAAAGATAAATAT
 AACCGAGTATGAATAAAATCCCATAATTTATTTATTTTCCACAGAAATATAGCAGCTTTCTGTGGACATTATGTCGCGCATATATGCTTAATGAGGTCATATACATCTGCC
 AACTCATATCGGGACATTACGCATACCGCAGGTAGACGACGAGGAGCAACAACTTTCTGATGCGCATTTTGGCAGGGCTTTATCTAAAGCGGCGATTAAATTCGAGTACATC
 AATCGGCTAATATGCAGAACAAACAAAAAAACTCTGGCCTTCCCCCTAGCAGCTAAGCTGTCCGCGACGCTGAGTGGCGAATAAATCTCGTCGGCATTTTGGCGGAATTAAGAGAT
 ACTTACGCAACAAAGATTGTGCCAATCAGTGTGGCTGGCAGGCAAGAGACAATGACCAGGATGTCGCCAATGCCCTTTTCGTTGGAGAGGGGGAATGTAAATTTTATTTCAA
 ATTATGGGGTTTCATCCAGGAGATTTCGGGCTCACCCACCAGATACAAATGAAGTAGTTCTGACCTGCGCATGCGAGGAGCCGAGACGACGGCGATCAGCAAGC
 TATTCCCGATGAGTCCCACCAGGAAGACGACGATGTAGGCCACGCAATGACCATGGACATGCGCAGCGAGTGACGGTAGAGTGGGTGCAATCCGAGTCCACGCTCCCGTC
 CAGCTGCTCCACCGGAGGGACATGCTGTTGTAGCACTTGGTGCCGCGAGCCGCGAGATCAAGCTGGTGGTTCGCGCGGCGATAGGTTACGAAGTTGGAATAGTAGGTCGGT
 GAAGTGGCGCTGCGGATGGCAGCAGATTGCCCGCAGGATCTCTGCGATGCGCTGTGTGGTGGCCCGCAAGTGGCTGACAGCAATGCGCCGGGAGCGAGCTGGG
 CCGTGGCCGTAGTGAGAACCGTGCCAGCAGAGTGTCCGTAAATGGCGCTGTTATTACAGTTTAAATGGCACTCGATGAGCCCAAGGAGCTGACAGCAGTCCCACCCTACCCT
 AGTGCCATTCAATTCGTTGAACTCAAGAGGTTGTGCAAAATCGCAGAAATATGCGTGAAATTTATGTAATCCCGCGGAGGCGCGCGCTCTCCCTATGCTCTCCGACGC
 ATTTTCCC CGCCCATGTCGGAATGGCATGGGTACACTAGTCGCTGACTGGGAACGTCGCCACTTGATGTGGCTTTATGCTTTTCGCTTGCGAATCCGCGGCTTGGCTGTG
 CCATCATCAAAATCATCGGCGCTCTGATTGGCGGGATGGCACTTTCACATGGCGCTCGCTTTCGCTATCATTCTGTAGACGTATCTCGAATCTTGAATCTGGTATCTCGC
 TTGTTTTAGAAAAGTCTATAAAAATGCCATCGTCACATTGGCATTTCGTTGGCTTGTGTAGGATATGCAGGAAGTTTCGTCGGCTTTGGCTTAGAGAACAGTTTAAATTAAC
 ACCCATGTCATCAATAATCTCGTTATCCGCGGGATGTTTGTGCTTTGCCCGAAGTTGCACATAAATCAGTTTCTAGAAACTCGCTTAGCGCCCAAGGCCCTTC
 CGTTTCGTTTATTTTATTTTGTGTTGAACTACCTACAAAAGTTTGTTCGGTTTCGGCTGCGAATCCGAGTGGCGGAATGCAATGATTGCTCGGGTCTGTCGACGTGG
 TTGACGTTGGCTTGGCTGGCATTACGAAAAGTCGGCGGCCCAAGGACCTCGTGTGCATATTTCAGATAAGCCAGATGCTGGGTGCTCGGATAAATGCAGCTAGAGGGGAAT
 GGGGATCGGCTTGGGCATTGGGCATTGGGCATTGGGCATCGGCATCGGCAAGGAGGCGAGAGCATCGGTCAGGACGATGTCACCGCCAGTGTGCAACACATCGC
 CGTCGTAATTAATTTTAAATCGCGCTCGCATTTGGCTGTGTTTGTGGCAATTGAAAGCCAGCAGCGTGCACTTGGACTCAAAAATGCAGACCAACATC
 GAACAAACGAGCAATTGAGCAACTCAATAGCAGCGCATAAAAATTCATTAGACAACTGCCAAGCGCAGCAGCACTCTTCGATTGAGTGCCAAAAATTCGCAGTAGC
 ATTTGAAACAATTTGTTAACAACGAGCCAGTGGCATTCTTTGTGCGGTTCCATGGGAGTGTCACCATCAGACAACTGGCGACTAAGGGCCCCCTGGCACAAATTCA
 (SEQ ID NO: 43)

Exon: 13111..12362
Exon: 12281..12156
Exon: 1649..1001
Start ATG: 13111 (Reverse strand: CAT)

Transcript No. : CT18916

ATGATGGGCAGCCAGCGGGCGGGATTCGCAAGCGAAAGCATAAAAGCCATACAAGTGGCGACGTTCCCGAGTACGACGACTAGTGTACCCATGCCATTCCGACCATTGGCGCCGG
 GGAATAATGGTTCGGGAGACAGATGAGGAAGCGCCGCCCTCGCGGGGATTACATAATTTACAGCATAATTTCCGTCGATTTCGAGAACCCTTTGAGATGTTTCAACGAATTTGAA
 TGGCACTAGTACGACCGTGGGACTGCTGTACGCTCTTGGGCTCATCGAGTGAATTAACATGAATACGCCGCTATTACGACAGACTCTCGTGGGACGTTCTCTACTACG
 GCCACGGCCACGGTGGCTCGGCGGCCAGTTCGCTGCTAGCCACCTTGGCGGCCACCACAACAGCATCTCGGAGAGGATCCCTGGCGGGCAAGTCGCTGGCCATCGCAGACG
 CCACCTCTATCCACCTACTATTCCAATTCGTGAACCATTCGCCCGCCAGCCAGGCTGATCTCGGGCGTCTGGCGCCACCAAGTCGTACAACGACAGATGCCCTCCGGTGGGA
 CGACGCTGGACGGGAGCGTGGACTTCGGATTGCAACCCACTTACCGGCTACTCGCTGGCCATGTCCATGGTCTACTGCGTGGCCATGATCGTCTCTCTGGTGGGACTCATCT
 GGGAAATAGCTTCGTGATCGCGTCGCTCTCGGGCTCCTCGCATCGGACGGTGACGAACACTACTTCATTGTGAATCTGGCCATTGGCGACATCCTGGTCAATTGTCTTCGCC
 TGGCCAGCAGCATGATTGGCAACATCTTTGGCGTAAGTATTCTTTAATCTCGGCAAGATCGCCAGGATTTATTTCGCCACTGGGGCGACCTTCATTGCCATTCGCTGGTGC
 CCTTGAACGAGATGACAAAGGACGCGTGGCCGCATGATATACTCCGTAATATCGGTAACTCGCGCTGGTGACCACTATCCTTGGCTGCTCTTCTCGACCTGGTGGCCGGCGA
 GGAGGTCTTCTCCGACGCCCTGGTCTCGGCCTACTCGACGCGCAGTTCCTCTGCCAGGAGGTGTGGCCGCCGGGACGGATGGGAACCTCTACTTCTCGTAGCCAACTGT
 GTGGCCCTGCTACCTGTGCTGCATCTGCTCCGTGATCAGCCTCTGCTACGTGCTCATCTGGATCGAGTCTCCACGAGTCCATCCCCGGCGAGTCCAAGAGCGCGAAATGGACC
 GCATCGACGAGAGGAGCAAGGTGAAGTCAATCAAGATGTGGTGGCGGTGGTCACTCTGTTTGTGTTTCTGCTGGCTGGCGCTCTAATGTCATCTTTGGCGGGATCAAAATCGG
 CTCGGACATCTCGAGGAGGAGTTGAGATCCTCAAGAGGTGATGCCGGTGGCCAGTGGCTGGGCTCTCGAACAGCTGCATCAACCCCACTCTACTACTAGTGAACAG
 AAGTACCGACGCGGATTCGGGGCATCATCAAGTCGCGCAGCTGCTGGGACGCGCTCAGGTTAGTTAAT
 (SEQ ID NO: 44)

Start ATG: 1 (Reverse strand: CAT)

MMAASGRIRKRKHSHSTSGDVPSTTTTSVMPPIPTMAPGKMAVETMEEAAALAGDYNNFTHNEVDLQNLLSFNELNGTSGSGGTAVSSLGSSSAIKLNNSAITDTLLGTVLTT
ATAFVAPASLLLATLATTTAAGSLAGSLAKSLAIADATSSYTNLNLSPATTSLLASAAATKSYNDSALRWEQDGSVDGFPDPLRHSLLMSMVCYVAYTVFVLGLI
GNSFVAVVLRAPMRMTVTNYFVILAIADIVFVLCPTLIGNFVRYKLVLSNARCRRDLFATGGHVCHLVAPEADDKATCPHNDNRHMGNAAGDHSLLAALLRPGARR
GGLLRPPGLGLLAAAVPLPGGVAPGHGWEPLLPASQPGGLLPAAHVPDHALLRAHLDQGLLHEVHPRRVQGRANGPHAAEEQEGHQDVGGRGHPVCFVLAALCHLCADQIR
LGLHLAGGVRDPQEGDAGGPPAGLLEQLHQPHLTLLSEQEVPTIRGHHQVAQLLRTPVQS*
(SEO ID NO: 45)

Name: Tachykinin receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384554

TGTGTCGACAGGCCACAGCAACCAATATGTTTCATTGTCTGCCCGGTTTGTGCGCTCAATTTATTTGTGATGTTGACTTTCACACAGTGACGTCAGTGAGAGCCACAGACAA
 TACCACATCATCTGCACCAATTAATAATGTTGTGGTGGCGAAAAACGCTCTGTGCGCCATTCTCAGTGGGGCGAAGTGAAAAGTGGAATTTCCAGCGCGTGCAGGAAATA
 GAATGAAATACGGGACATCTTGAGTGGTGTCCTTTATCTGCTTCACACATAAATATGAATAAATTAGCTTAAACCAAAATGACTACAAAGTATAGCTCTATTAA
 CTCCTATGGTCTACTTATGCTTAATGGGTTTGGTTAAGATGCTTGGGCATGTTATCCACCATTACGAGTGGCTATATACACGGATGAGTGGAAATTTAAATGGCCCTTA
 TAGCTGATAGGAGCTGAAAAGAAATGCATTTAAATTTCTGATAAATGGCCCGGATTCACATACAAGAAATCAGCTGTTTAAAGAAATATAACTAAACCGAGCTACTAA
 TGGATTGAACCTAGCTCTGAGTGGTGGACAAAATACTTAATAGTTTTCGAACACTATTTCATTAGGATACCATATACTTAATTAATTTTCCAGTTTAAAGTGAAT
 TCGAAAAATGCTCAATTAATGTTTCAGGTAAAGTTTGTGAAATATAAAACCTTTGTACCAATACAAATATAATGGGTTCTTAGTATAAATTTAAATAAAATCAAAT
 CTGGAATAGATTCAATACCAATTTGACACTTAGATTGTTTAAATTTAGAGTTAATGAATACCGGTATAAATGTATCTTAACATAAGATTGCGAGTAGAAATAGGTTTG
 AAATTCGTAGCGTAGACTTGGCGTACTACGACTAGAAACATACATTTGGCTTACCATGCCCTGGTATAACTAICGAACCTATTGTCTAAAGCCCTCTGGTGGCTTCCACAGGA
 TGTCGGGTCTCCCTCTTAGAGAAATACCCGCTCGCTAAAGACCTGGGACCATGAATCGAGAACCGGGAATCTTTCCGCGTGGTGGGCGGACCATCATACCTCCGACATG
 CGCCGACGCTGAACCACTTTGGTGGGACGTTCCGAGATTACACACGGCGGCTGCTGTCTTTGGTGGACCTTCGGGAAACCGCAACGAGTGCTCAGTGTCTTCATCGAAT
 CGCAACTGTGTGTAAGCCCGACGCCAACCGTGGGACGCGGTTTGGCTGCGAGTTTGTGGGATAGCTGCAGCAGCATGTGAACCACTTAAAGCGGGAAGCGCATCGGA
 AATATAATAAAAAATAAAATATCAAGTAGTAAGTAATTAAGCATAGAAATATGTAAGATTTTAACTAAGAACTTAAACATACTTAATGTGCGGAAGCACTGGCATGA
 GAGAGGACATAGATATTAGTGATTTCGCCCGGAGTTACAGCGTGCCAGACTTTGGATGAAGGTGGCGATGGCAATGTCGCTCTGTAGTGTGGAATCTGGCCAAAGACCTGCA
 AGCAGATCGGAAGTGATATACCGGACCAACGAGCATGATGCAACAAACATGATGGCTCAATGTCTTACCTTTGACCTTGGCCGTGGGAATATGCCCTCGAGCTGG
 CCTCTCTGGAGGTTGAGCTTCCGCAACACGACGCTGTCTGTAATAATATAATGAACAAATAAATATAATTTTGGGAAGCTCCGATTTGGGAATATGCAATGACGCAAGAG

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TATCTTACCCTGGGTACAAAAATGGAACCTTTGCCCAAAATCGTCTTTACGATGATCGCATAGCAGGCAGATATGATACAGCGAGGAATGGCAATAGAGTGGCCGACACC
AGGCTCATGTACACCTGCCAGGCGATCGGTGAACCCCAACTCAATCCAGCATTCGCGGATGTCTCTTGGATGAGCTTCTCTCGTACAAAACCAGGATGGGAAGCGAAAAACAGC
CCGAGATGAGCCATGCGCCGCGACCGAGGTGACGGGCTTTTTCCCTAAAAGCATAAAATTAATATTAATGAAGAAATATATTTCTAAATTAACCTAAATTTGTATA
TTTTTTTGGTTAATCCCGAACTAAATTTATCCAAGCATGTAGCAGAAGCCAACACAGCGCTCATCTTTGGCTAACACAACTTTGGCAATTGAAAACTTTTCATACGAC
GACTTAACCTACACGATTTGAGAAGTTCTAGGGGTGTGTGATGGCATCGTATCTGTCGATGCTCATGCGCCACAGCAGCTAGGTGGCCAGCGATGTGACGACAGCTGCGA
GAAGCGATGGCCTTGACGGCAGGTTGCTTCCGCCCGACGAAATCGTATGCGGCATATGATGTCGTGGAGACGCTTGAGCAGTCCACGCACAGATGTGAAATATATGGA
CTGGGCATTAGAAGCAAGGGAAGAGGTCAGTCCCAATCCCAAACTCACTTGCCAAATGCCAGCTGTTTAAATGAAGTAGTTCATCCGCGACTTGCATTCTTGTGATGAACA
TCACGAACAGAACAGTGAATTTGCCAGAAGCATGACGGTGAACAGGATCCAGAGCAACAGCAAACTGTTCGGCTTGGAAGAGTGCAAGAACAAGTGTGGTCCATTTTCAT
TAAACGCCATCAAGTTCAAGTTCAATGTAACAAAGTGTTAAGCAGCGAAATCAGCCATACCAACTCAATTTAATGCCATCAAGCACACACACACACAGCGCTTGTATTGGCT
TTTAAATAGCCATAAAATGATAGAGCTATGAATCAAGCTGGTTGGTGGCGCGGCGAGGATGTGCCGGCGGAAGTTTGCTTCTTCCAGTTTCAGTACTAACAAAGCATAGGAC
AACACAGCATCTCTGCATCAAAACCCGTGGACGCTCACCACACACCCAGCATGTATGGAAGACAGACAGCACCCGTTTATTCAGCAGGAAATTTGCTGCCAAGATGAAGGA
CTGCGACTCGAAAAATATTTCTTTCGACTTTAAAGGCGATTTCACAGTCCAAATGCGTGGCTGGTATTTAATTTATAGAACAATATACCTCCAGCGATTGCAAAATCAATT
CATAGCACTCAAACTCAAAATTTGTACAATTTTATATAGAAACTAGTAGATAAAGAAATATGAACCCCTTATTACTTTTAACTTTTAACTGAACTGTTATTTATTA
ATATTTTAAACAGTGTGAAGTGCAGATATGACAGTGAAGTCTGCTAGTTCTAGCTTATTTGCGAAGAAATGAATATTTCTTATACGCGCCGATGCGATTTATATAGC
AGAATATCGCATATATGCTAATATGACTAATGATCTCTTATGATATCTACAAAGTTTTTTGCTGCAAAATTAATTTTGGCTCAAGGCACTCCCCCTATGCCACGCTCAT
TGAAGTTAAGTTGAATTTAGGCTCTGGGGTCTTGACGCTGACGCGCAACCCGATGTTGATGCGGACTTACTTAAGGGCAGGCCAAGTAACTTTGCCGAAAAAAAAGCT
GAAATCTGACAGGCTTCGTCTTGGAGTCTGCGCTCAGACTGAAGACGCCCCACCTTATTTACAACTTTTTGTCCATACCAAACTTCAGATATTCACCTTTTTG
(SEQ ID NO: 46)

Exon: 2688..2514
Exon: 2450..2253
Exon: 2061..1801
Exon: 1716..1431
Exon: 1339..1111
Exon: 1042..1001
Start ATC: 2688 (Reverse strand: CAT)

Transcript No. : CT19191
ATGAAATGCTGACCACATCTTCTGCTCTTGGCACTCTTCCAGACCGCAACAGTTTGTCTGCTCTGATCTGTTTCACCGTCATCGTTCTGGGCAATTTCAGCTGTTCTGTTCTGTTGA
TGTTTCATCAACGAAGATCGCAAGTCGGGAGTACTACTTCATTAACAGCTGGCATTTGGAGATCTGTGGCTGGGACTGCTAACGTCCTCACCGGACATATATGGCGCAT
CACGATTTCTGTTGGCGGGCAGGCAACCTGGCCTGCAAGGCCATCCGCTTCTCGCAGGTCTCGGTCACATACTCGTCCACCTACGTGCTGGTGGCCATGAGCATCGACAGATAC
GATGCCATCACACACCCCATGAACCTTCTCAAAGTCGTGGAAAGACGCCGTCACCTGGTGGCTGGCGCATGGCTCATCTCGGCGTGTGTTTTCGCTTCCCATCCGTTGTTGT
ACGAGGAGAACGCTCATCCAAGGACATCCGCAATCTGGATTGAGTTGGGTTCCCGGATCGCTTGGCAGGTTGTCATAGCCTGGTGTGGCGGACTCTATTGGCATCTCTGCT
GCTGATCATATCTCGCTGCTATCGCATCATCGTAAAGACGATTTGGGCAAGGGTTCCATTTTGTATCCACAGGAACGTGCTGGTTTTGGAGAGCTGCACCTGCCAGGAGGGCC
AGCTCGAGGGGCATTATTCCACGGGCAAAAGGTCAAAACGGTCAAGATGACATTGACCATCGTGTTTGTGTTTCATCATCTGCTGGTGGCGGTATATCATCTTCGATCTGCTGC
AGGCTGTTTGGCCAGATCCACATCACAGACCAACATTTGCCATCGCCACTTCATCCAAAGTCTGGCACCGCTGAACCTCGGCGCGAATCCATATCTATTGCTCTTCTCTC
ATCGCAGGTCTTCGCGACATTAAGTCGCTTTCGCGCTTTAAAGTGTTTCACATGCTGCTCGAAGTCAATCGCACAACACTCGAGCAAAAACCGCTGCCACACGGTTGGTGTG
CGGCTTCAACAACAGTTGGGATTTCGATGAGGACACTGACCACCTCGTTGACGGTTTCCGGAAGGTCACCAACAAGACGAACGCCCGTGTGTTAATCTCGGAACGTCCCACCA
AGGTGGTTACCGTGCACGCCATGTCTGGAGCGACGCGGAGTTTCTCTAAAGGGGAACACGGACATCCTGTGA
(SEQ ID NO: 47)

Start ATG: 1 (Reverse strand: CAT)

MKCDHTLFFALFQTEQFAVLWILFTVIVLGNASAVLFVFMFINKNRKSRMNYFIKQLALADLCVGLLNVLDTDIWRITISWRAGNLACKAIRFSQVCVTSSTYVLVAMSIDRY
 DAITHPMNFESKSWRRARHLVAGAWLISALFSLPLVLVYEELKIQGHPCWCTELGSPIAWQVYMSLVSATLFAIPALISACYAIIVKTIWAKGSIFVPTERAGFGAAPARRA
 SSRGIIIPRAKVYTKVMTLTIVFVFIICWSPYIIFDLLQVFGQIIPHSQTNIAIATFIQSLAPLNSAANPLIYCLFSSQVFRTLRPPFPFKWFTCCCKSYRNNSSQNRCHTVGR
 RLHNSCDSMRLTTLTSLTVSRSTNKTNARVVICERPTKVVTVPAMSERRGVSLKGNTDIL*
 (SEQ ID NO: 48)

Name: Vasopressin receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384294

TTCCAATGCGATTTAAGGTAATTGACTATAAGTACCACCGAAGCTCTCTGGACAGTCTCCCAAGTGGAAAAAAAACCTGGATTTATTTCTTACGAAAACCTGCGATAAC
CAATTTGTATATATGTTCTATTCAGGAGCGTGC CGCATTTTGTAGTTTGC GAAATGAGTTGTTATGTTTCTGAGTCATTGTCGAGCGGGTTTTGGTAGCGGGGATTAATCT
CGTGTCATCATCTTGGTCCAGCGTTATTTCTATTGCTTTTAATTTTCACATCAAAACAGCAATAATAAACCGTGCCCTATCTTACGGACATATCTCCCAAGCGTCGTCT
CGCTCTAATTTGGATTTTTCCTCTCTCTCTCGCTCTTTTCTTTCGACGTGTGCGGCCGACGGTGGGAATTTCACTACGCTCTGTGTTGTCTATTTCACAGCAGCATCTGGCCT
ATAAAAAATGCAATTTTATGTTTAAAGAAATAGTGC GCGAGTCCGCTGTGTTGCACATTTGGAAATTTGTCGAGAAGCGAAGAAATGTGCTTGGAAATGGAAAAACA
CGAGCAGCATAGCAGCAGCAGCATATAACATTGAGCCAGATGTGAATACAGAGCAAAAGTGAAGCGCGCAATTGGCAATTTGCAAAAAGAAACAGATTATACAAATTT
GTGTTGCTGCGCAGCTGAACCTAGGACAACAGCAGCAACAGCAGCAACCAACCACTAGTGTATAAAGACCCGGCTGGCAACAGCTGCAGCGGCCATTGCAAGCAACAT
AATTTCAAAGTCTGGAGCTGGGAATTCGCAAAATCAGCGAGTGGCAGCAACACAGCAACTGCAGCAACCTCTGCGCTCTCCGGCATCGCTCTTTCGCTCGCCG
TTCTCGCGCTGGCGTTTCTCGCTTTCGCTCGCACCCCGCTGTGGAGCAGTCGCGTATATTGGAGCGGTGGGGCGAACCTCTACGACAGATTTGCTTGGAAACGCTCGGAG
TGAAGATCGTGCTCGTCCGATCGGTGCGCGTGATAAATACACGGCGAACAAAAATTGAGTCAAGAGTCAGACGGTCGCAAGACAAATTGACAAGACACGAAAAATCATAATAA
CAAGAAGGGTAAAGCAACTGCTTGAGACAAAAAAGAAATGTTTTTACAATTTCTCGCAGCTATAATGACAAGCTTCGCAAGAAACAAACCCGATACAGATCGAACA
AAAAAAGGGATTTCAACAAAAACAAAAATTTTGGCTCGAAATGTGAAGACGACGAGACCGGATATACTACCGAGTGAATTTGTCTACATACAGACGACAAACA
AAAAACGAAAAAGCCAAATTCATAAATAAAGGCCATAAGTGACTGTGAACCTGTAAGATGATGTGTGTTAATCGAAAAACAGCATAAAGAAATGGCCATAGACCTCCTGATCTGGC
CCTGCTGCTCGTCTCGTTCGTATCAACCTGCTGGCGCTGTGTGCTTTTGGATTAACGCTCTGAGCTTGAGGACCACCGCAATCGCTTTACGATCAACCTGCTGGCCATCA
CTCATTTGTTGCTGATCTTTGGCCCCCATTTGTTCTTGGAGTCCGGGAAGTCTCGGAGGCTACCTTCCAACCGCGAGACCTGGAGTTCTTCTGGAAGCCGGAA
ACCTACAGGTGATCATCTCGTCCGAATGGCCAGCTGGTGGAACAGGACGGCGTGGTGTGCGAAGGAACATCAGCGAAACCGCGGATACGGTGGAGACCTTCTCAAGTGC
TGCCACCTATCTGTGCGAGTTGACCATCGATGAGCCGCGAGATGTTGCTTTGTCAATCAGCGAAACCGGACACACGAGAGAATCTCTCGGCCTTTGAGAGGTTACCCAG
GAGGCTCCCATTGTTGCCGCCCGTCCAATCTGAGATGTGGTCCATAGACATGACTGCTGCGCTGGGAGCTCTGGCAGTACTGCTGCTGTTGGGGACACCTGGTGTGCGGTT
CGGATCCCTCGCTACCAAGCTGCGATTTCGGCGGTGAAACCTGGATCTTCATAGACCTACCTGGTGGTGGGACATCTTTTGAGGCCCTTTCGCGCTCCGAGTTT
GGACTTTGAGCGGACGCGCTTTCAGCAGACAGACGCCGCTGGCAGTCACTGATCTCAACATCAGCAACCAATAGTATTTTCGAGTGGTGTATGCGCAGCTGTC TACTTC
ATCTGTGATCTCTCTCTGCTTTCGCTTCTGCTGTGCGGAAATGTGATCGGAGGATCTTCAAGTGGGCGCGGAAATGGGCTCGCGATCGCGGACATGGCTCTTGC CGCTGC
TGCAGAGCGCCTTGAATCTGACGCTTGGCAGCGGCGGCTCAAGCCCAACAGTTCTCCAATAGCTTGTGTGTTTCACTCGGCATTCATATCTCGGCGAGTTCGATCGCGG

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CAACAGCAGCTTGGGCTTGGGCGGTCTGCAAAATGCAAAATGATCAGCGCCAAACAGCCGAGGAGTTGCGCCAGTTGTCTCCGTCGGGACTCGGCCGCGAAAGTACTTTTGGCCC
ACCATCTCGGATGATGGTGGCTCAGATCGGAAAGTGGAGCCGGAGTTCAACTGATGCCGTACAGGAGCACTCACTGTCAGATAGAAATCAGAACATTATGCTTACTCTGC
AAACAGCCAGTGGTGAATAAGAGGAACACTCCGCCAGACAGCTGCCCTGTGGGCACTTCTTCGAGGATCTGAGGGAGACTAATCGTCTCAGGGCATAACGCCAGGT
GCACAGTTCCGCCAATCTCCACAAGTACACGGAACCTGCCGAGGACTCGCTCAGTGAGGAGTGTGGTTCTCCGCATCTTTGGGTCACGCCACGCGGAGCAGCAGCAGCAA
TTGCACCTGCAACATCAGCAGCAACAGCATCATCACCACCAGCAACATCATCCGCACTTCAGTTCCGCCGCCACCAGCAACACGGCCAGCCCTCCAGATACCCGCCATTTC
ATGCCCTCCCCAAGGCTCTGAGTTACATGAGCTCCTTGCACACCGCTTTGAGCAACGCCAGTTTCGCTATTCAAATACCGTGAGGAGTCGAGAGCAGCCCGCATCAGCATCTCT
GGTGGTGGTGTGTTTGTGGTCTCGTACCTGCCCTTCGGACTGCTGGTACTCTGCAATCGCGACTGTCGGCTGCTAACTTTGGCGGATCCTCCAGCTGGCCATCTTCATG
ATCCTGCTGGCCAAATCTCAGTTCCGCATTTCATCTTCGCCCTACAGGAACAAGAGGGTGAGGCGAGGAGTCAAGCGGCTGTTTGGTCTGGATTCTGCTCGGGCTGCAGCGCA
ACTGCAGCAGCAGTGTAAAGACCAATGGCACCAGGTCCTGCTGCAGTGGTGCACAAATTCAGCGCAACAGCAGCAAGTTGTCGCGAGTACAGCAGCAACAGCTGCAAGTA
TCTCACCCTCCAGAGTTCCTGGTCTAGTCAAGTCCGCTGCATACGACCTTGACGTTAAGGCCCAACAGCAGCTGCAGCACCATCATCAATTTCCGGTGGCGCCAGGGCTCA
GCGGACTCCGATGAGCAGCAGCCGCGCAACACCACCCTACTGTGGCAGTTGCTCCACC GCCACAGCGCAGCGCCCAAGCTGCAGAGGGCATCACCATCGTTGAGC
ACATAGCTATAACACCAACCATGCCCAAAAGTTCCAGAATCGCGCGCCAGGCTCTTCGACATGTTCTTCGCGAGCTCCAGAACTGCAGGCGGGCTGCCAGGCCAGAG
TTTGCCCACTGAGGTCTAAACCGGGGTTATGATAACTATACATATAGGAAGGTAAACAACAACTAACTTGTCTGAAGAATTCTAAGAACAGTATTCGTAATTTGATA
GTCTAGCAATTAACAAGACAACCTGTTTGAATTTGAGTTTAAAGACCAGATCATGCTGTCTAGAACTTCTCATGCCAAAAACAACCAAAAGAAAGTCAAACTAGAA
AGTTTGAATGCTTTAAATTAACAAAAAAGAAAAAAGAAAAAACAACAAAGTACCTTTTAAAAATAGAAAACTGTGTACAGTCAAAATTAACAAAGATTGAG
AGTGATGGGAGTTATTAATCTTAGGTTGGATATCAAACTGAAAGTGAACCAAGCCAAAGCAAGTGCATCAATCAATGTACCTTAAGCAACTTGCATATGAACATTAA
TGTTTTTACTCAGATGAAATCAGGAAATAGCAAGACAGCAGCTTATTCGAACCTCCCAACTAGTTTATTATTATTGAAACAATTGTAATATTTATTAGACATC
CAGCGCGCAAGCTTTAACTCCAAAAACATTCTGGCTAAACAAGAGAGCGCATACATTTCGATTTCATAGTTAATTAAAGTCAATGTGCAAGCATATGTGTATTCTAGAGT
TTTGGCCAGTGCACAGTTGGTTTTCAGTTGGCCAGAAATGTCAGTCAGCAGTATGTTAGCCTTTGGTTAGCCAACGGCTATTGGGCCATTAAATCAATTTGCTTTTAGAA
CACCGCGGGCATTGCGTGCAGCTAAATACCTCATTAGTTATAATTTGAATGCGGATGATGGCGCAATAGGTCAGAGGTCGGAGGCCATCGGCACGGCCCATGCATATAG
CATGAAAAATGTTGATGCCCTGGCATTTGATCATTGCTTTACTATGCTGCAGGCCAGAAAGCTTTATACCTTAAGTTCTCTCTGTTGATTTCCTTTGGCTGCACAA
AAAGCAACAAAGCAACACACTCACACAACACAAAAAACCTCGAATTTATATTTCTAAAGTTTATATTTTATAAGAGTGTGTCATAGGAAATATCTATAAATATTCT
ATGCATGTGTG
(SEQ ID NO: 49)

Exon: 1001..4051
Start ATG: 1431

Transcript No.: CT19320

CTTCGGAGTGAAGATCGTCTGTCGATCGGTGCGCGTGATAAATACACGGCGAACAATAATGAGTCAAGAGTCAGACGGTCGCAAGACAATTGACAAGACACGAAAAATC
ATAATAAACACAAGTGGAAAAGCACTGCTTGAGAGCAAAAAAGAAATGTTTTTACAAATCTCGCAGTAAATTAGCAAGCGTTTCGCAAGAAACAAACCCGATACAATAC
TGAAACAAAAAAGGGTATTCAACAAAAAACCAAAATATTTGGCTGCAAAATGTGAAGAGAGCAGAGACAGCGGATATACTACCGAGTGAAATTTGTGTATAGCAGAG
CCAAAAACAAACGAAAGCCAAATTAATAAAAGGCCATAAGTACTGTGAACCTTAAAGTATGTTGGTAAATCGAAAAACAGCATAAAGAAATGGCCATAGACCTCTCTG
ATCCTGGCCCTGCTGCTGCTCTGTTCTGATCAACCTGCTGGCGCTGTGTGCTTTTGATAACCGCTGGCTTGAGGACCACCGCCAAATCGCTTTACGATCAACCTGCTGG
CCATCAATCTCATTGGTGTCTGATCTTGCCGCCCACTTTGTTCTCTGGGACTGCCGGGAAAGTCTGCGGAGGCATCCACTTCCAAACGCCGAGACCTGGAGTTCTTCTCGAA
CGCGGGAACCATCACTGTCGCGAGTTGACCATCGATGAGCGCGGAGATGGTGGCTTTGTCATCAGGAAACGAGACACACGAAGAGAATCTCTCGGCTTTTGAGAGCT
AAGTGAATGCCACCTACTGTCGCGAGTTGACCATCGATGAGCGCGGAGATGGTGGCTTTGTCATCAGGAAACGAGACACACGAAGAGAATCTCTCGGCTTTTGAGAGCT
TACCACGAGGAGCTCCCAATTTGGCGCCCGTCCAACTGAGATGTTGGTCCATAGACATGACTGCTGCGCTGGGAGCTCTGGCAGTACTGCTCGTGGTGGGGACACCTGGTG
TGCGGTTACGGATCCCTCGCTGCTGCTGCTGCTTTCGGCGGTGAAACCTGGATCTTCATAGCACTCACCTGGGTTGGTGGGCGATACCTTTGGAGCCCTTTCCGCCCTTC
CGAGTTTGGACTTTGAGGCGGACGCCCTGTTACGACAGCAACCGCGCTGGCAGTCACTTCAACATCAGCAGCACCATAAGTATTTTCGGAGTGGTGTACGCCAGTG
TCTACTTCTGCTGATCATCTCTCTGCCCTTCGGCTTCGTTGTCGGGAATGACTGAGGAGTATTCAGTGAGGCGCGCGGAAATGGGCTGCCGATGCGCCAGAAATGGCTCTTC
GCCGCTGCTGAGAGCGGCTTGAATCTGACGGCTGGCCAGCAGCGCGCTCAAGCCAAACAGTTCCTCAATAGCTTGTGTGTTTCATCGGCATTCATATCTCCGCGCAGTTTCG
CATGGCGCAACAGCAGCTTTGGCTTTGGCGGCTGCAAAATGCAAAATGATCAGCGCAACAGCGGAGGAGTTCGCCCACTGTTCTCGCTGGGACTCGGCCGCGAAAGTAC
TTTTGCCCAACCATCTCGGATGATGGTGGCTCAGATGCGGAAATGGAGCGGAGTTCAGCTGATGCCCGTACAGGAGCACTCACTGTCAGATAGAAATCAGAACTATTGCT
TACTCTGCAAAACAGCCAGTGGTGAATAAAGAGGAACACTCTCCGCAAGACAGCTGCCCTGTTGGGCACTTCTTCGAGGATCTGAGGAGACTAATCGCTGCGAGGGCATA
CGCCAGGTGCACAGTTCCGCCAATCTCCACAAGTACACGGAACCTGCCAGGACTCGCTCAGTGAGGAGTGTGGTTCTCCGCATCTTTTGGGTACGCCACGCGGCGAGCAGC
AGCAGCAATTCACCTGCAACATCAGCAGCAACAGCATCATCACCACCAGCAACATCATCCGCACTTCAGTTCCGCCGCGCCACAGCAACACGCGCCACGCCCTCCAGATACC
GCGCATTCAGCTCCCTCCGCTGCTGAGTCTTGGGACACCGCTTTGAGCAACGCCAGTTTCGCTATTCAAATACACCTGAGGAGTGCAGAGCAGCCCGCATC
AGCATCCTGGTGGTGGTGTGTTTGTGGTCTGCTACCTGCCCTTCGGACTGCTGGTACTCTGCAATCGCGACTGTCGGCTGCTAACTTTGGCGGATCCTCCAGCTGGCCA
TCTTCATGATCTGCTGGCAATCTCAGTTCCGCAATTCATCTTCGCCCTACAGGAACAAGAGGGTGAGGCGAGGAGTCAAGCGGCTGTTTGGTCTGGATTCTGCTCGGGCT
CGAGCGCAACTGCAGCAGCAGTGTAAAGACCAATGGCACCGCAGGCTCCTGCAAGTCCGCGCGGAGGCTTCGACATTCGAGCGCAACAGCAGCAAGTTTTCGAGTACAGCAACAGC
TGCAAGTATCTCACCCCCAGAGTTCCCTGGTCACTGAGTCCGCTGATACGACCTTGACGTTAAGGCCCAACAGCAGCTGCAGCACCATCATCAATTTGGTGGCGCCA
GGGGCTCAGCGGACTCCGATGAGCAGCCACCGCAACACCACCACCTACTGTGGCAGTTGCTCCACCGCCACAGCAGCGCGCCCAAGCTGCAGAGGGGCATCACCAT
CGTTGAGCACATAGCTATAACACCAACCATGCCCAAAAGTTCCGAATCGCGCGCGGAGGCTTCGACATGTTCTTCGCGAGCTCCAAGAACTGCAGGCGGGCTGCCAG
AGCCAGAGTTTGGCCACTGAGGTCTAAACGGGGGTTATGATAACTATACATATAGGAAGGTAAACAACAACTAACTTGTCTGCTGAAGAATTTCAAGAACAGTATTGTA
ATTTGTAAGTCTAGCAATTAACAAAGCAACCTGTTGAATTTGAGTTTAAAGACCAGATCATGCTGTCTAGAAGCTTCTCATGCCAAAAACAACCAACGAAGAAAGTCA
AAGTGAAGTTTGAAGTCTTAAAT
(SEQ ID NO: 50)

Start ATG: 431

MAIDLILALLLVSLINLLALCAFWITPGLRTTANRFTINLLAINLIGCCILAPTFLGLPGKSAEASTSNAETLEFFSKPGNHQVILRRNGQLVEQDGVVVRNISENGD
TVETFKNATYCRELTIEDRGDDGFTVETETHEENLSAFESLPTAEPLPPVQLRCWIDMTAALGALAVLLVVGDTWCAVTDPLRYHSRISGVKTIWIFIALTWVVGILF
GALSARVLDFAEDALFSRQRRLAVTYFNISSTNSIFGVVYASVYFIVIIILPFGFCVMYWRIFSEARGNGLRMRQNGSSPLLQSLANLNTAGQQAAQANQFNSLCLVHRHS
ISSASHHGNSLGLGGLQMQIDQRQPRSSPSLRRDSAAKVLPTISDDGSDAESGAGVQMLPFQEHSLSDRNQIMLTQATSGEIKRNSYARQLPLLTSSQDLRET
NRLQGITRQVSPNLRHYTELRLQSSSEECGPHLLGHAGRQSSQQLHLHQHQQHHHQHHPHFSRPHQHQHGHALQIPAIHASPALYSMLLRHLSNASSLFXRYEE
SRAARISILVVVMFVVSYPFLGLLVLLQSRLSAANFGSSQLAIFMILLANLSPPFIAYRNKRVRRGVKRLFLDSSSLQRNCSVVKNTGTAGPAASGALQQRNSKLS
QYSSNSCKYLTQSSVLSVQPVHTTTLRBNSSCSTIIFEGARGSDSDSEPPATPPPTVAVAPPTRPQRPRKLRGITTVEHIAITPTMPQKFNRRARLFDMFFRSSKK
LQAGCQSQSLPTEV*
(SEQ ID NO: 51)

Name: G-protein coupled receptor-like
Classification: G_protein_linked_receptor

FIGURE SHEET 25

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ATTCTATTGTCGTAGGCGTGGCAAAGGTTTTTGGTAAATCGATAGAAATGTACAAGAGAATGTAATTAATATGAAAAGGTAAACAAAAATTTGCAAAAAGTGTGGGCATG
GCAGTTTGGTGGCTGGTGTGGCGTAAGGGCATATCGGTATTTTCTTAATTTCCAGCTATATGATGCAGTCGCCTAATTTTGTATCAACTATACTAAATTTGTTATTTGTC
AGAAACAACGACGATGAATTTATAGTCATTTATCTTTTTATATTCTTTATCTAAAGTATTATGATATAGAAATAAAAAACAGTACGAGTCTCTTAAGCGGACGGAA
TAATGTCCTCGAGATCTGGACCTTCATTAAGACAGACGAACATTAGGACAAGGCCAGGTCTACTCAGCTAGGGATCCTGATCAATGTATATGTATTTACTTTAAAGGGGCGTA
AAGTTGTTATTTTACATGTTACATACCTTTCAAGCGCAATCTAGTAACCCCTTTAATTTTACAAGTACAAGATATAAAAAGTTGAATGGTGAAGCGGAGAAATTTAATTTGTTAAT
ATAATATTATTTCTTTGTTAATCCAGAGCTTGGCAAGCTGTGCACACAATTTTTTTTTTTTTTTTGAAGAATCAATGATTAATTTGGCCAAATTTAGAGTGTAGATTTCTT
GGAAATCTCAATGAATTTTATGGTCAAGTTTCCAAGAAGTGAATGCTTAAACAAACAACAAAAACAAACACACAAAAGCAAAACAAATTTATGCAGAAATTTACGGATGTATTT
ATATGTATGTATGTACATACATATAACAAGAAATAAATATGGTGGGTCTGTGGCTCTTGATGAATTTTCAAATTTAAGCTGCAATACATGTACATTTAAACGCTTTCTGCT
ACGCTCTCAAATAAAAAACAATTTTAGGATTAATAAAAAACAGATATGGTGCTTCTATATCCCATAGAAGAAATATTCATATGATTTTTGTTTGGCAGCAAT
GCGAATAGTTATTGGAATCGTTACCGCATTTCTTTTGCTGTTATTGCAAACTCAAATGCCGAAATTTCCCGGTTGCGACTTCTTCGACACCGTAGATATTTCAAAGCGCCA
AGATTCTCGAAGCGAGTCGACTCTACGAGGCTGCTGATCCCGCCCATTTGACAGCTGAATATGATCACAAGCTCTGGCGACGATTCGAAGGAGAAGGTGGCGAGCT
ACGTACGAGGATGTGGCTGCCACCTCAGGCCATGCAATTCGGTTTTGTTGCCCTTGAACACAAAGATGCAAAAGACGAGTGTACGGCGACATTCGGGAGGCGACTGAA
CAAGCAGCATCCCTTCGTGAACGTGACGCTCAGCGACGGGTCCGGTGGTCAGGAGACACTTCAAGGAGGATCTGATCGTGCAGTCGGATCTGGCCAAGCCCGGATGTCGCCGG
AGTCTACTTTCTGAATCAGCAAGCTGCCGGCAATGAATTCAGCTGTTTGGAGTTACTTTCTGCTGTTTCTGTCACCTTGAAGATGTAAGCTTTTGAATCTCTGTAGAA
CGGCTCTCTCTGCGTCACTGGGACAAGTGAATCTGACAAAGCCGGAGTACTGCTGCAGTACTTTCATTAAGGATAGTAGTATCCGAATTCGACCCCACTCTGCTCC
ACTTTCTGCTGAGCACTCAGAAGCTGGAAGACCGTTGGTAAGCTGAATTCCTGCTTGACTTTAAGGTTTTTAGTTAATGTCCATTTTACAGCGATAGTGATATCCTTGA
TAGTCATAATCTTGAAAGTACAGCGTGATCACTCTAATTTGAGAAAGCTCGCAAACTGCACGGAAAGTCTTAACTGCTCATCTGGCCTCTGTTCTGGGATATTTCTCTGTA
GGTCCCTCAAGCTGACGAATATCTCTCCGGCTTCTGCGTTACAGCAGGTTTGGCTTGGCGATACAAGTAAAGGACGACCAATTTCAATTTTCGATATCATGATCCGGT
TCCTGGGCTACTTCTCCGTCTATCGCCGGTCTTTTGGCTTTCTGTCATCAGTCTTACCCTCTGGAATTCATTTAGCGGCAACTCCAGCTGGTTGAACGTTTCTTGCCACA
GAATCGGTTCTTCTTCAATTTATACGCTCGGGCATGGCGTCTCTGCACTGCAATCACTATATAGCTGACAGGCTGGTAAAGACGAGAAGTTGAGACCGCGGCTG
GGCGTCGGCAAAATTTGTGATCTACAGTAAGTTACTATTCCCTATAAACAATTCGGGTGCTGAAATCACTCGTTTTTTTTTAAACCAAGCTGGGATATGACTGTATGATCT
ACTTCTATGGCCCGATGCTACTGATAATGTTTTCAACATAACAATGTTTGCTGACGGCTTTTCGTATAATGAAGTGAAGAAGGAAGCACAAACTTTACTCAACAGCA
AAAAACAACCAATAGGCTCAATTCCGACAGCAGACGCTGAATATTTTACCATATTTTATACAGTCTTAATCCAAAGTTTAAATTTTAAATTTATTTCTATGGCTTTTAAAG
GTACGCCCTATGCTCGCACTCTTTATCATCATGGGCTGCTTGGAGCTTGAAATCACTCGTTTTTGTGAGCAAAATCAGGCTTGGCCCAAGGCTTTATGGTGGC
TGATTACTCAATTTGGTCACAGGACACCGTCAATTTCTGCTATTTGTTCTGAGGCCAGCACACTGAAGCTACTGAAGGAACGGTAAGTTATCGGTTCTTCCATAGGAGGT
GCAATTTAACTGGTTTATCATCTCGGCTGGGTCAGGAGTAAAGGTTGGGAGGATGAAGCCGGCCAGCGATGAACATATCTCACTTCAGAAATCGAAATTTGACCCAGCTGT
TTTTTAATATATGGGTAATGTGAAATACAGTTTATAGTTTCATTTGGTGTGTGAAACCTCTATTTTGAATCACTAACAGGACGCTAATGATCGTGACCAAAAGGCTGCTT
TGCCTGCTCATTTGCTCACAATATCGTATGCTCATATGGGCTACGAATATTTGCTACTTTATCACTTTCAAGTATCTTGATTTCTGCACGGTGTCTAAGCATATCTAAGTT
TTAATTTAAAGTGTGAATGTAAATGCTCTTTAAGTTATTAACTTTGTGCAATTAATATCTTATCTCAATCCATCTAGAGCTGGCCATCAAGCGGATACGAAGCACTACAT
GGGATATGTACGAATCTTATAGGCATGAATCTCGGCATTTATATACCTGGTCAATTTATTTAGTATGAGTGAGGATAAAACTCAGAAATACTTTTGCATCAGCACTTAT
GTATGCTCACTCGTGAATAAATTTCTGAATAATTCGTACTGTTTTTACTGCCCAACTTAATTCAAAGAACTACAATGTATTATATTATGACTTTTTATGACAATGACTAA
CAAAAAATAAAAAATTTCTGAATTTTATAACAACCTTTAAAAATGTTATTAACCTTAGGATAAATAAACAACAGTTTAAATTTTACAAGCATCTGTTAAAAACAATAA
TTACTTGAATATAAATTTAAAAAAAATGAATATGGTTTACATTAACGCTATGCAAAAAATCAAAAAGCTACAGACTGACAGATAGAGAAAAATTTCAAAAAAAGTA
TCTAGTCGAGTTCTCTGCACTATAAGATACTCATTTCTCAGCTAGTTCTTTCATTTAAAAATTTCTCTTCTGTTTCCGATAGTAATTCGCAAGAAAAATAATAAATACACA
AATTAATGTCATTTTGGAAATTTATGGGCGTACTGATTGGTTCGGTTAGTGGACGTGTAATTTATGAAGCAAGCGTGACACTACAACAGAAATATATATATATAT
(SEQ ID NO: 52)

Exon: 1001..1507
Exon: 1567..1703
Exon: 1756..1951
Exon: 2014..2268
Exon: 2329..2500
Exon: 2578..2772
Exon: 2835..2919
Start ATG: 1007

CAGGAAATGCGAATAGTATTATGGATCGTTCACCGCATTTCTTTTGCTGTATTGCAAAACTCAAATGCCGAAATCCCGGTTGCGACTTCTTCGACACCGTAGATATTTCAA
AAGCGCCAGAGTATTTCGACACGGATCGTCACTTCAGAAAGCTTGCTGATCCCGCCCAATTGACAGCTGAATATGATCAAGAGCTCTGGCCGACGATTCGAGGAGGAAGGT
GGGAGCAGCAGTCAGGAGATGTGGCTGCCACCTCAGGGCATCATTCGGTTTTGTTGCCCACTACCAAAGATGACAAAGAGCAAGTGTACGGCGGACATGCTCGAGAGC
GAGCTGAACAAGCAGCATCCCTTCGTGAACGTGACGCTCAGCGACGGGTCGGTGGTCAGGAGACACTTC AAGGAGGATCTGATCTGTGAGTCGGATCTGGCCAAGCCCGGAT
TGCCCCGGATGCTACTTCTGATACACGAACTCGCGGGACATGAATTCAGCTCTGTTTGAGAACGGCTCTCTTCTGGCTCACTGGGACAAAGTGGAACTGAGCAAGCCGGGAGT
ACTGGCTCCAGCATCTTCTATTAAAGATAGATAGTATCCGAATGCACCCCACCTCTGCTCACTTTCTGCTGAGCCATCAGAAATTAATGCTCATTTTCACAGCGGATAGTGA
TATCCTTGATATGCATAATCCTAACGATCAGCGTGTACCTCTATGTTGAGAAGCTTCGCAACCTGCACGGAAAGTGTTTCACTCTGCTACTTGGCCCTCTTGTTCTCGGGATA
TTTCTCTCTGGTCTCAACGCTGTGGGAATATCTCTCGGCTTCTGCGCTTACAGAGGGTTCTCTGGGCTACTCTCTCGCATCGCCGGCTTCTTTGGCTTTCTGCTCATAGT
CTTACCTCTTGGAAATCATTTAGAGCGCAACTCCAGCTGGTGAGCCGTTTCTGCCCAGAAATCGGCTCTCTTTCTCAAAATTTATAGCCCTGGGCA TGGCGGCTGTCTCTGCA
CTGCAATCACCTATATAGCTGACCAGGTGGTTAAGAACGAGAAGTTGAGACCCCGCGTGGCGCTCGGCCAAAATTTGTTGGATCTACACTGGGGATATGACTGTCTATGATCTA
CTCTCTATGGCCCGATGCTACTGATATATTGTTTCAACATAACAAATGTTTCTCTGACGCTTTTTCGATATTAAGTAAGTAAGGAAGCAACAAATCTTACTTCAAGCAGCAA
AAACAAACCAATAGGCTCAATTGCGACAGCAGACTTACGCCCTTATCTCTGCGACTTTTATCATATGGGCTGTCTGGAGCTTGGAAATAATCTGTTTCTTTGTAGGCA
AAATCAGGCTTGGGCCAAGGCTTTATGTTGGCTGATTACTTCAATTGGTCACAGGGCACCCTCATATTTCTGCTATTGTTCTGAGGCCCAGCACACTGAAGCTACTGAA
GGAACGGATTAAAGGTGGGAGGGATGAAGCCGGCGCCAGCGATGAACATATCTCACTTCAGAAATCGAAAATTTAGCCCAAGTGTTTTTTAA
(SEQ ID NO: 53)

Start ATG: 7

MRIVIGSFTAFLLLLLQNSNAEIPGCDFFDVTDISKAPRFSNGSYLYEGLLIAPHLTAEYDYKLLADDSKEKVASHVRGCACHLRPCIRFCCPQYQMKQSKCYGDMSEDEL
NKHDFVFNVTLSDGSVVRRHFKEDLIQVSDLAKPGCPRMYFLNHLELPGNEFTLLFENGSLLRHWDKVELSKPGVLRPASFI*
(SEQ ID NO: 54)

Name: mth-like 3
Classification: G protein linked receptor

Celera Sequence No. : 142000013384666

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AATTAAATGCATTTTGGAAAATATGGGCGTGACTGATTGGTCGGTTAGTGGACGTGTCATATTATGAAGCAAACTGGGACACTACAACTGAAATATATATATATATATAT
 ATTTTCATTGCCCTTAAGTGTGAAAACCGCTTTAAATTTGAGAAGTACTTTTAAGAGATATTCGAGGATATTCGGTGACACCGGCACACTTATCGCATGGCCATGGGAATAT
 TTAGATTTTGTAGAATAGGGTAATATGTGAACGTAAACCTTCGATTGTATAAAGCATTCAACTCTTAATATGATATATGCTGTACGGTTCACCGGCTATTAACATTTCGCGTTTAT
 TTACTTAAATTTATCTTTCTATGTACATGCCAGAACTGATAGAAATGGCACGGTTACCGCCGATAGTCGCTGAGTATGGGGTATCTTATAGTCGAGCACATTCTCGTTCTCC
 CTGACATTCGCATATATCGGTTGAGTGTGTCGAAATTTGGGATCAGTCACTTAAACCTCTGGTCCGATAAACATGTTTTTGTGTAGTCGTCCATTGTGACCTTCGGCTAT
 AGTATTTATATCAAGTTTTCACGATTTTGGATCGTTAAGCATAGAAGGCCACTCGAGTACGATTTGGACGTTTTTATACAATTTCTGCTCTCTTAGACGAAATAAATAT
 GTGGCTGCCCTGTGTTTTTAAAATGAACATTTGCTTACGCTTCGTAGAGAATGCTTAGTCATAACAGATGGATTTGTTGAATTTATTTTTAACAGTTGATAAGTTATACTTTAT
 AAAAAAAAAAAAAAACCTTACTCAGAAATGCTATAGCACATGACTCAATTTTTGTTCATTAAGTATTTAGATCGTTTCAGTTTGGCGAGGTTATTTTGGCTACTTTAT
 TTCTAATTTGTGAACACGTAACCTTGGTTTTAAAACCGTATATACTACAGTAAGTATCATTTTGAATTCAGTTGCTCTATCATATTAATTCCTTCTTGACGAAATACGGAT
 ATTTGTAATTCGACGTTCTTTTCTTGCTGATGCCAAAATCAAATGCCGAAATTTCCCGGTTGCGACTTCTTCGACACCGTAGATATTTCAAAGCGCCAGATTCTCGAACGGGA
 TCGTACCTCTACGAAGGCTGCTGATCCCCGCCATTTTGACAGCTGAATATGACATCAAGAGCTCTGGCCGACGATTGCAAGGAGAAGGTTGGCGACACCGTACGAGGATGTG
 CCTGCCACCTCAGGCCATGTCATCGGTTTTGCTTCCCGCAGTACCAAAGATGCAAAAAGTAGCAAAAGAGCAAGTCTATGGCGACATGTCGGGAGCAGAGTGAACAGCAGATCCCTTT
 CGTGACGCTGACGCTCAGCGACGGGTGGTGTGAGGAGACATTTCAAGGAGGATCTGATCGTGCAGTGGATTTGGCCAGGCGCGATGCCCCGATGTACTTTTGAAT
 CACGAACCTGCCGGGCAATGAATTCACCTGTTTGAGGTTACTTTCTGTGTTCTGTCACTCTGAAGATCTGTAACCTTTTGTGAATCTCTGTAGAACGGCTCTCTCTCGCG
 TCACCTGGGACAAAGTGAACCTGAGCAAGCGGGATGACTGCTGCCAGCATCTTTCAATTAAGGATAGATAGTATCCGAATTCGACCCCACTTCTGTCTACTTCTGTCAGCAC
 TCCAGAACCTGGAAGACCGTTGGTAAGCTGAATCCCTGCTTGACTTTAAGGTTTTAGTAAATGTCCTTTCACAGCGATAGTGATATCTTTGATATGCATAATCTCAAC
 GATACGGCTGTACCTCTATGTCGAGAAGCTTCGCAACCTGCAACGGAAGTGTTTCACTCTGCTACTTGGCCTCTTTGTTCTTGGGCTATTTCTCTGGTCTCAACGCTGTGG
 AATACTCTCTCCGGCTCTCGGCTACACGAGGTTTGGTTGCGGATATCAAGATAAAGGACGCAATTTACATTTTCGATATCATGATCAGGTTCTCTGGGCTACTTCTC
 CGTACGGCCGGCTCTTTTGGCTTTCTGTGATTTGGAATCCACTTGGCGATTAAGTTTGTAGTCTAGGCTCTAAGCTTTTGACCGGATTTGCTGCCAAGCAAAATCCGTTCCGTGCA
 TACAACCTTATACGCCCTGGGTTATACCGTATCATGACTGCAATCACTTATACAGCTGACAGGTTGGTTAAGAACAGAGAAGTTGAGAACCCCGGCTGGGCTCGGCAAAAT
 TTGGATCTACAGTAAGTTACTATTTCTATAATTAACAATACCGTGAACCTCGTTTTGTTTAAACCAACAGCTGGCAGATGATGCTATGCTACTTCTATGCCCCAGTGT
 GTTCTAATCGCTTTCAACATAATAATAGTTTGTCTCTTTGCGGATTTACATATAAATCAAGAAAAATGTGAAGCGCTGTGCCAAGCAACAAACAAACCAAGATATA
 AATGACCAACAAATGTAATTTATATATTATATAGATAATACTATAATGTCAATATGGCACTTTTGTGAATCTTTTATAGGTTTGCATATTTCTGCGGACTTTTCATCTTAA
 TGGGTTTGTGCTGGAGCTTTGAGATATCTCTTTTGTATAACCAACAGCAAGCTTGGGCTAGGGCTTTAATGGTGGCTGACTATCTTTAATTCGCTCCAGGTTACCATCAT
 ATCTCGGCTTTTATTTTAAAGCTAGATTTCAAACTTATAATAGCAGGAGTAAAGATATTAACAATAGCTGAAATTTGACCGTCTCATGCTATGAACATCTTT
 TGATTTAGGGGACGTCAAACCTCCAGGAAGTCAACATAATTCGAGATCAAAGCAGCTCGATATAATCAACTCATACGGCTTGTGAAGGATCAATTCGGGATCCAAACG
 CTTACTCTCAAAAAATGGAATTAATAAGATGCAACCTCTCTGTTATTTATGATCACTATTTAATTTCTATGGAGCAAAAAGTTTAAAAAAATATTTGTTTATCAAGCAA
 TAAATATATGCTTGGAAAGATTAAGAAGAAATATATAAAGGTTACAAAAGACGAAAGATGTGGGCAATACAGCAAAATATTTTCGGATCATCACTTTTGTGTAAGA
 CAAATGTAGAAATATTTCTGCGACCGTTCTTCTTCAAACCGAGGTTTGAAGTCAACACTGTGGCCACCACTGCGCAAGTACCAAACGATACAGAAATATTTCTAATCGAT
 TAGCACTCGGGGGTACATCGATGTTTCTCGGCTTCAAAATCAATTTCTATGTTGTTTTTGTGCTGGGCTTTGACGTTTGTGCGCTTTCCTCCGTTGGCAATATTAATATAACG
 TAAGCTATTTTATTAACAACGGAATTCAGCGCACACGACGACCACTAGTAGCATACTGTAAGCTGAAGCGAGGGGCAAAAATTAATTCGGTTTGGCGGCGAAGAG
 TTGATGACGTCGCATACGCCGTCTCTAGGCGTAAAAAGCAAGCAAGCAAAACCAACGGAAGCGAAACGCTGTAACCGCGTAGAAGCGATAAACGCGACTCAAATACG
 CAGCAGATAAAATCAATACCGGAGAGAGAAAAGTACGCGGAAATATTGTTTCATA
 (SEQ ID NO: 55)

Exon: 1001..1492
Exon: 1552..1702
Exon: 1759..1935
Exon: 1998..2252
Exon: 2627..2640
Start ATG: 1001

Transcript No. : CT2035I

ATGCGGATATTTGTTAATGTCAGTCTCTTTTCTGTGTGATGCCAAAATCAAATGCCGAAATTCGCCGTTGGCAGCTTCTTGCAGACCGGTAGATATTTTCAAAGGCCCAAGATTCT
CCAAAGCAGTGCCTCTCTACGAAAGGCTTGTCTGATCTCCCGCCCAATTTGACAGCTGAAGATGATCACAAGCTCTCTGCCCAAGCATTCGAAGAGGAGAAGGTGGCGAGCCAGTACG
AGGATGTGCTGCTGCCACCTCAGGCCAGTCATGCATGGTGTGTTTGTGCCCCCAATGTACAGTACCAAAATGTGCAAAAGAGCAAGCTGCTATGGCCAGCATGTCCGAGGAGCTGAGCTAACAGAC
GATCCCTTCGTGAACGTGACGCTCAGCGACGGGTCCGTTGTTGAGGACAGCACTTCACAGGAGGATTCGTATGTGTCAGCTCGGATCTGCGCAAGCCCGGATGTCCCGGGATGTACT
TTTTGGAATCCAGCAACTGCCGGGCAATGAATCTACTCTTGTGAGAACCGCTCTCTCTCCGCTACTCATGGGACAAAGTGGAACTGACGCAAGCGGGAGTACTGCTGCCAGCATCT
TTCATTTAAGGATAGTAGTATCCGAATGTGACCCCACTTCTGTGCCACTTTGCTGTCGAGCACTCCAGAAGCTGGGAAGACCGTTGCGATAGTGATATCCTTGATATGCATAATC
CTAACGATCAGCGGTGATACCTCTATGTGCGAAGAGTCTTGCAGCAAGCTCAGCGGAAAGTGTTCATCTGCTATGCTGGCCTCTTTGTTCTGGGGCTATTCTTCTGGTCTCCACAGT
TGTCGAAATATCTCTCCGGCTCTTCGGTTACAGCAGGGTTCTTGGGCTACTTCTCCGCTATGGCCCGCTTCTTTTGGCTTCTGTGATTTGGAATCTCACTTCGCGGATTAAGTT
TAGTCTAGCCTCTAACTGTTTGCACCGATTGCTGCCAGAAAAATCCGTTCCGTGTCATACAACTTATACGCCCTGGGATATACCGCTAATCATGACTGCAATCACCATAACAGCT
GACCAGGTGGTTAAGAACGAGAAGTTGAGACCCCGGTGGGGCTCGGCAAAAATTTGTTGGATCTACACAAGCTTGGGCTAG
(SEQ ID NO: 56)

Start ATG: 1

MRILLIAVLFLMPSKNAEIPGCDFFDVIDISKAPRFSNGSYLYEGLLIPAHLTAEYDYKLLADDSKEKVASHVRGCACHLRPCIRFCCPQYQKMQSKCYGDMSEDELNKH
DPFVNVTLSDGSVVRRHFKEDLTVQSDLAKPGCPRNYFLNHELPGNEFTLFFENGSLLRHWDKVELSKREYCVQHLSFKDDSIIRIAPHFCPLSSEHSRTWKTVATVISLICII
LTISVYLYVEKLRNLHGKCFICYLASLFLGLYFFVLNVWVYSSGFCVTAGFLGYFSVMAAFWLSVIGIHLRIKESLASNCLHRLLPENPFAYNLYAWGIPLIMTAITYTA
DQVVKNEKLRPRVGVGKNCWITYTSLG*
(SEQ ID NO: 57)

Name: mth-like 4
Classification: G protein linked receptor

Celera Sequence No. : 1420000I3384544

GATTGCCGTGGTCGTGCGCCGCGGGAGAAATGCCCTTTTCGTGTACGCATTACTAGGCCAAAAGGTTGGACGCGCCTTGCTTTATTAGGCCAATAATTTACGCTGTCGAGC
GGCCATTAGAGCGGCATCCCGCTGGCCAAAGTTGGTCTATAAAAGCCGGCGGCTGCAAGTGGCTTTTCCATCCACAGTGAAGAGACACTCGAGTCGGCGGCAGAGATGCT
GCCGCACCTGCTCGCCAGAGAACAGAGTCTTGCCTGTCATCTGCTCATCTGCTCTGCTGTGACGATTTGAAACTCTGCAGGCCACACCGGATAGCTCTTGTCGCC
GCCACGGATATGATGATTGTAAGGCATTGATAATATCATTAATTCGCTCTAATGGATTCCATTTACCATCAGATATAGCATCTTGTCGGCAGTGTGATCTGAGTACGAGC
CCCGCTATCCATGTGCTCCAGTATCTGATGTTGTGCCCAAGCCGATTCCCTCGTCTGCGCTGGTGTGCTGTGCAGTTATATCCAGTGGGGTGGATTATATCTTGAAG
TATAAATATAAATTTGATATAATCTGATGACTTAGGTTTTCGGGCAGTAAGATCTGGCAAAATGGAGCGTTCTCGCATGTGCTGCCAGGAGTCTGGTGAGCGGGAGGCAGCCG
TCTCGCTATTCTGCTCGCCAAAGTGAGCCCGCGGAGGCTAAATTAAGAAGATCTGCAGCAAGGCGCATTTGGAGTGCATGTGTCGGCCATGCATCTCATTAGGAGTCTGG
CATCATACCCAGGAAATTCGCGGTATTCGCGACGAGGGTCCACTCAAACTTCTCCGGCGCATTGCTGTGCATAAGATTCCCCCATCAGTTTAGCATCATCAATCATGC

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Exon: 6503..6410
Exon: 6016..5754
Exon: 5577..5096
Exon: 5034..4854
Exon: 4798..4580
Exon: 4524..4229
Exon: 4168..3805
Exon: 3742..3632
Exon: 3565..3341
Exon: 2916..2571
Exon: 2508..2403
Exon: 2345..1879
Exon: 1817..1001
Start ATG: 6002 (Reverse strand: CAT)

29/89

AAAAGTTCCTCGTTGGCTGAGATTGATGAAAAAAGTGTTCCTCAATGGCCATCGGTAGTGAATAAAAGCCAGTTCTGGGACTAAGCAAGCTGCCACCGCGGTCAACATGT
 TCCGGCCAAAGTGGTTTCCATTCGCCACCGCTGCTGCTCTGCTTCTGGAGCAGCCGCTGTGGCAGGACCGCAAGAGATCGGACGCTACATACGGGGATTCTCCCGTAT
 CGGGAGTGGCGTGAATACCTCTACACCGGTGGGGCGTATTGCCCAGTGAAGCTGCCGTTGGGTCACGTTAATGAGCATGGAAAGATATCGGCCAACTACAGGCTGCAC
 ATGTGGTGAACGACACTCAGTGCAATCTGCTCTGTGGGCGTAAAGTCCTTCTTCGATATGATGCATTCCGGTCCCCATAAAGTGATGCTCTCGGGCGTGGCTGACCCCATGT
 TGACCGATCCCATAGCCAGGCCAGCAAGCACTGGCCCTCACCCAGCTCAGCTACCGGGACCCCATCCCATGTTCCACCAAGGATGCGTTTCCGAATTCTTTCCGCTGGT
 ACCCTCGGAAGTGCCTTTAATGCGGCCGCACTGCCCTTGCTGAAGGAGTCAATTGGACAGAGTGGGCATGTCTACAGGATAGGCCACGCTATTTCGCTGCCCCAAAT
 CACATGGTGGCTGACCTGGATGCCATGGAGTGCAGGTGGTGGAAACGCAGAGCTTCGTCAACGATGTGGCTGAATCATTTGAAGAACTGCGCGGAGAAGGACGCTGAGGATGT
 TTCTGGGCAACTTTAAACGAGCACTTGCACGCAAGGCATTCTGTGAGGCTTATAAATTGGATATGTATGGCAGAGCCTATCAATGGCTGATCATGGCTACCTATTCCACGGAA
 TTGTTGGAATGTCACGCAAGCAGCGAGTGCAGTGTGGAGGAGTGCCTGACGCTCTGGAAGGTGCCATTCTAGTGGATCTTTGCCCTTTGCCACAGGTGGTGACATCACA
 GTGGCTGGCAATTACTGCTGATGAGTATGCTGTGGAGTAGCAGAGCTCGGAGGCATGAATATCCCGCTTTCAGGCTATACCTACGATGGTATCTGGGCAAGCTGCCCTGT
 CCATTCTAGTATGTGGCCGAAAAAGCAGAGGATCTGCTAACACATTTTGATTAATCCGCTGAAGGACGTGGAGAGTGCTTCTTGGAGCTGACTAGTAACTATCTTCAGGG
 TGTGACGGAGCCGTGCGTTTCTACAAACAGCGGCAAGGCCAACATCCTGATCAATCAGTTTCAGCTGGGACAAATGGAAAAGATCGGGGAATACCACTCACAGAAGTCA
 CATCTGGATTAAAGCTGTGGAAAACCACTCAAAATGGTGGGGAACACTCCTCCAAAGATCGCACTTTGATCTACATCAGCAGCAGTCAAGGTCAATCCCAACCATATATAATTG
 TATCCGGCTAGTGGCTTCGTGGCTATGGAGTGATTATTTGCCACAGTTTTCCTGGCCTTAACATTAAGTATCGCAATCAAGATACATCAAGATGTCAGGTCCCATTTGAAACAA
 TCTGATCAATTTGGGCTGTATGATGACCTATTTGAGCATCATTTTCTGGGCTCGATACCACATTAAGTAGTGTGGCAGCTTTTCCCTATATCTGCACAGCTCGAGCCTGG
 ATCTTGTATGGCTGATTCAAGTCAAGTTTGGAGCCATGTCTCGAAGAGCTGGCGGGTGTCATTCGATATCCACGGATCTGAAGCTCAATAAGGAAGGTCTCAAGGACTATC
 AATTGTTTATGTTGTGGGCTGCTTTGGCCATGTATATGCCATATAACCACTCGGACAGTGGCATGCCCTTTACCGCGAACTAAACAGATTTGAACCTCTGCATCA
 CGAGAAATATTGATGATGCTTGGTGATCCCCGAAAACGAGTACTGCCAGTCTGAGCAGATGACCATATTCTGTAGCATTAATTTAGCCTACAAGGAGTCTGTTGTGGTTTT
 GGGCGCTTTTGGCCGTGGAACTCGCATGTTCTTACGCCGCTCGAAGCATTTCAAGATCATTTGGTTTCTCCGTTTATAAGCTGTTCATCTTTGCTCGGCCGGAGCG
 CTATATCCCTGGTGCTATCGATCGACAAGGATTTAGTTTGTGCTTACTCGATGCTTTTATCATTTTGTATCGACAGCCACTTTGTTGTTGGTGTCTGTCAGCAAAATGGT
 GGAGCTGAAGCGGAATCCCCAGGGCGTGGTGGACAACGCGTTAGGGCCACGTTGAGACCATGTCCAAAAACGGAGCCGCGGGATTCCTCGGTGTGCGAACTGGAGCAACGA
 TTGGCGAGATGTAAAGACAACCAACTCGCGATTCCGAAAGCGGCTGTAGGAGAAGGAGACAGAGCTGCAGGCCCTTAATCCGCAAGCTGGGAGCCGCGAGGACGCAATGGATCG
 ATGGGTTGACCTGCACAGGTGCAATGCCGCTGCTGTAGGACGCTGAGCGCCATCTGAGCAGTGAACGATGACATTTGAGGCTCTAGCTCGCCGGTGCAGAGATGCCGACGAC
 CACAGAAGCTTACCGAGATTACGCTCGTGATAGTGACCTGCATCATGTGGAGATGGATAACTCCTTTGTGTGGTGACGTTACAGTGATGGCGCATCGCTTCTCTCC
 AAAAAAGAAAAGCAATCGATTGTAGAGCACCATCGCATGCCCTGCTCCCACTATGATCGGCCCATCCAGCAGCACTTCAGCAGCAACTACAGCAACATCAGCAGATGC
 AGCAGACGACCTGCAGCAGCAGCAACACAGCAGATGCAACAGACAAGCAGCAGCAGCAGCAGCATCATCTGCCATCTGGAGAAGAGAAACTCGGTGTCCGCTCAGACGGA
 TGATAATATAGGCGACATCACCAGTACGGCGGCAAGCGGAGCGGAGGAGACTGCTCAGATCGGGAGAGGCGTCAATCGACGCCCTCCAGGCTACTACGACATGCGGACG
 CAGACGCCACCCCGCCGGCCAAAGTACAGAGCTCGCACCGGAACTCTCCACCAATCTCCACATCTCCCATCTGCAATCTGGAATGTGAGCAACATGTGCTCACATCAAAGCCGAGTA
 CTCGGCTGTGATTAAAGCTCCCACTGCCCTCGGACATCGCGCACAGCATGGGCTCCGCTCTGAAGTCCAATTTCTGTGGTTTACAGAGTGAACCTCTGGGACACGCGACAC
 GCTGTCGCGACCGCAAGCAGCGCAAGTCCGCGGCAACTACGCCAGCTCGCGACGCTGTGGCAAGATCATGCGCGCCAGGATGACCTATGACCCGCAACACCATCTCGCC
 ATTCAGCGGCTCGCTCTCGAGAAGCAACCGCCAAACATCGGCAAAACCGCAAGGCGACCGTTTGGCAGAGCAGACGACGAGGGAAGGGAACGAGATCGCCGCGCCA
 ACAGTCAAGCGTGGCTCCAGCCGCGTAAGGTACGCCGAGCTCTAACATCCAGCACGCCGCCACCACCAAGTTCGCCCAATGTGGCGCCCGATAAGCAGCGGAGCAGGCA
 CGCGCGCAAGCAGGATAGCAGCATCTACGGCCCGCAGCAGGAGCAGGAACTGCGAGGGGAGACGCAATTTGGCCACTCTCCGGAATCTCTCCAGGAGAGAGTCCC
 AACTATCGGGCGGCGAGTGGCTGGGCGAGCTGTCGCAATATATCCATCAATCGCATATGCTCGAGTACTGTAGCGGCGCGAGCATGTGATTCATTTGCGGTAGTTAA
 ACTCCTATTGTTAGATCCAGTTGCAGTGTAAAGATTATCTTTGTTTCTTTGGAGTTAGCATCTACATACATACCCTGTGAAGCAGGACACACCATTTAGTGAAGTACTAGT
 AGCAATGACATGAGATAGTGGGAAATAAATTCACATTTAAATTAATCATATA

(SEQ ID NO: 59)

MFRPSWFFPASLLFLLLWSTACGRATAKRSVDVYIAGFFPYGDGVENSYTGKRVMPKVKLALGHVNEHGKILANLYRLHMWVNDTQCNAAVGVKSFDDMMHSGPNKVMFLFGAACT
HVTDPIDAKASKHHWLTQSLYADTHPMFTKDAFPNFRFVVPSENAENAPRLALLKEFNWTRVGTVYQNEPRYSPLPHNMVADLDAEMEVEVETQS FVNDVAESLKKLREKDVIR
ILGNFNEHFHAKCEAYLKMCDMYGRAYQWILMATYSFHTTDDWNVTDQSCSEVEEITALEGAILVDLLPLTSGDITVAGITADEYLEVVDTRLGRTEYSRFGHYTDGIWAAA
LAIQYVAEKREDLLTHFDYRVKDWESVFEALRNTSFHNGTGPVRFYNNERKANILINQFQLGQMEKIGEYHSQKSHLDLSLKGKPVWGWKGTPEKDRTLIYIEHSQNPNTIY
IVSASAVGIVIATVFLAFNIIKYRNOYKRMSEPLNNLIIIGVCMNTYLSIIFLGLDITLSSVAFFPYICTARAWILMAGFSLPFGAMFSKTRWHRVHSIFTDLKLNNKVIKD
YQFAMVVGVLDAIDIAITITQWADPFYRETKQLEPLHHENIDDLVLPENECYQSEHMTI FVSI IYAYKGLVLGFALFAMEWTRHVSIPALNDSKHIGFSVNVVFTCLAG
AAISLVLSDRKDLVFVLLSFFIIFCTTATCLCLVFPVKLVRKNPQGVVLDKVRVATLRPMKSNRRGRDSSVCELEQRLLRVLKNTNCRFKALMEKENELQALIRKLGPPEARUK
IDGVTCTGGSSNQVLEPLINDIVRLSAPVVRMPSTTEVNTSDVSTSTHVENDNSFVSVQSTVMAPSLPPKKKKQSIIEVHSHASAPATMMQIQQQLQHQHQQHQ
MQOQHLOQQQHQQMQQQQQQQHHHRLEKRNVSQAQTDNIGSITSTAGKRSGGDCSNMRERQSTASRHVDSGQQTPTARPKYSSHNSNSTSTQSELSNMCPSKP
STPAVIKTPTASDHRRSTSMGSALKSNFVVSQDLWDHTT LSHAKQRQSPRNYASPORCAEHGGHGMTYDPNPTSP IQRSVSEKNRNKHKRPKPKQKVCQSETDSEERERDP
PNSQPCQVQPRKVRSSNSTQHAHHHSS PNVAPDKRNSRQRGQDSSITYGASSETELEGETAILPIFLKLLTEKS PNYRGRSAVGGQSCPNI SIKCDIVEYIL*
(SEQ ID NO: 60)

Celera Sequence No. : 142000013383823

GTTCCTTAAACGTGATCAAAATATAACATAGAGCTACTACTAAGCAGCGAGCAGCGCTGCGATATATCAACAAAGTCCAGCTGGCAACATGTTGCCGCGCCTTGGGTGAGTT
GTGCCCTATGCTCTTTGGCCAGATTTGGCGGCGCTTGGCGCGCGCTGGTGTCACAAAAAGCGCCGAGCTTCCGCGTACGTGCTTCGCGAAGATTGCGAGGCGGCACACA
CATCGGGAAGCCGAACCTCCATACGGCTTGCGGCACATCAATCCCGCATTAAGCCCAATTTTGTGCGTGAAGTACGAAGTGTGTGTACTGCTGCAAGTCAAGTCAAGT
CCCCATTGCTAGTGCTATTCCGGGATCACAAACGAGCATCTCTTTCGGCCTACGATTACGCGCTTCTACCATTACTTCGCGCCAGGCGCTGCCGCTGGCCCAAGGAGCGTGC
GATCCATCTGAATCCGACACAGAGATCTCCGCTGCGGTGACGGGGTCCACATCAACAGGGACACCGGTGATCTTCTCAACTACCGGGGCTGGAATTGGATCTCGGACTG
GATTTGGATCTTAATCTGGACATGGAGCTAGCCACTACTCCAGACGACAGCACTTGGCGCGAGCTGTAAACCTGACGACACCGGGAACAGAGCGCTGGTAGGGTCAGTG
CCGATGTGCCCATCTGGGTGGTGCCGTGCTATTCGGCCATCTCTCTGTGCGCCGTGGTGGGAACCTGCTGTGTGCTGTACCGTCTGACCTCCAGAACTTCGATCGCATGCGCACCAT
CAGCAAGCTTTTCTGCTCAACCTGGCCATCTCGGCATCTCTGCTCGGCTATTCTGCTATCCGCTGGTCCGCACTCTGTAAGGCATTTTCATATTCCGGCAGAGTGT
CTCTGCAAGCTCATCCAGTTTGTCTCAGGGTGAAGTGGTGAAGAGTAGGCGAGTTCAATTTATGTCTTGTGCGTGTCCAAATTTAAAGGTTATCCCAAACTCTCAACTGTGA
ATTCCGCATTTATTAGTCCCTATCCCGTGACCGAGCGCCATAATCGCTTGTCAAATGGCCACGAACAGCAGCACTCGAGCGGATTCGCCACAGATCGTGTGGCCAAAC
ATTACAGGAGCTGTAAAGAGATGACAGCTCGCTGCTTCGACATGTATGCTGATTCGCCGAAATGGAGTGTGCGCTCACAGGAGCGCACTGCCCTGGCCAACTGTATGGATC
GATTCTAGGACTCGGTGAAGTGGTGTGCGAGCGAGTACTTCGCTGCGCGGTGCGCCATCGACAAATCGTTTGTCCGCTTGACCGCTCTCCGATCACCACAGCAT
CGCATCCGCATCCATGCCAAATCCGACAGCAGAAATGAATCTGAATCCGCATCTAGGGCCTCCAATGACGAAAAGGTCAAAATATGGTTAAAGTCTAGACAGATAGCCACGCT
GTGGAATTTTATCTTCAAGTTCGAAACCAAGAGGCTGTTCGCTGTAGCCCGAGATCCCGAGTACTGTGGTGATGCCCACTTGTCACATGTGTCACATCCCAAGATTTTCGA
GGATGATTTCATACCTTCGAGACCTTTGGGAAGGGGCAAGGGGGTCTGTACTGCGGAATGAAAGTGGATTTTCTGACGGCTGGCCATCGATGATTTCTGCTATTTGGTTG

FIGURE SHEET 31

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AAAGATCAATCCCAAAGTAAAGGTATTCTGATCACGGTGCAGTGAAAGGTTTCAATGAATCAAGAGCATTTGACAGCAAGCTGATTTTACAGGGTACACAGGGAAGTCC
CACTCTCAAAGGATTCCCATCATTTCAAAGTGGTGTGACAGCGAGCTTATGGAGTGGCAGACACTCCGAGGTCCAAACTAAATTAAGAAACCAATCGGGTGCCAAAT
CCAATTTAAGTAGAGCAAGTAGCTGGAATGCCAATCGGATATATTCGGCGGGATTAGCATTTGTG
(SEQ ID NO: 61)

Exon: 89..105
Exon: 176..214
Exon: 1025..1252
Exon: 12710..12910
Exon: 12965..13303
Exon: 17202..17499
Exon: 17579..17996
Start ATG: 89

Transcript No. : CT21155

ATGTTGCCGCGCTGTGCGCCGACGCTTCCGCTCAGTGCTTCGCGAAGATTGCGAGGCGCGACACATCGCGGAACCCGAACCTCCATACGGCTGTGCGACACTCAATCCC
GACCTAAGCCCAATTTTTTGTTCGCTGAAGTTGACGAAGTGTGTTGACTGCTGCAAGTGCAAGTCCCGGATTGCTAGTGTATTTCCGGGATCACAACAGGACATCCTTCTT
CGGCCCTCAGGATTGACGCTTCTACCACTACCTCGCCGAGCGCTGCGCTGCGCAAGGAGCTGCGATCCACCTGAATGCCAGCAACGAGATCTCCGCTGTCCGTGACGGG
GTCACCATCACAGGGACACCCGGTGATCTTCTCAACTACAGCGGGCTGGAATTTGGATCTCGGACTGGATTGGATCTTAATCTGGACATGGACCTAGCCACTACTCCCAGCA
GCAGCACTTGGCGCCAGCTGTAACTGTCAGGACACCCGGGAACAGGAGCGTGGTGAGGGTCACTGCGGATGTGCCATCTGGGTGGTGGCTGCTATTCGGCCATCCTTCT
GTGCGCCGTGGTTCGGAACCTGCTGGTGGTCTAACGCTGCTCCAGAACTGCTGCGATGCGCACCATCACGAACGTTTTTCTGCTCAACTGGCCATCTCGGACATCTGCTC
GGCGTATTTCTGATGCGCGGTGACCTTGGTTCGGCACTCTGCTAAGGCATTTTCAATTCGCGGAGCTGCTCTGCAAGCTCATCCAGTTTGTCTCAGGTCCTATCCCCGTGACCG
AGCCCCATATCGCTTGTCAAATGGCCACGAAACAGCAGACCTCGAGCGGATTGCGGAGGATCGTGTGGCCAACTTCAAGAGCTGATAAAGAAGATGACAGCTCGCT
GCTTCGAGTATGTATCGCTATCGCCGAATGGAGTTGCGCTCCACGGAGCGGACTGCTGGCCAACTGTATGGATCGATTCTGGACTCGGTTCACTGCTCGGTGGCC
GTTTCTCTGAGCACTGCTGGCCATCTTTCGCGAGCGCTACTACGCCATTTGCCACCCGCTGAGGTGCGGCACCTGGCAGACGATCAACACGCCCAACAGATCATCGCCA
TCATCTGGTGGGAGCTCTGGTGTGATGACGCCATCGCCGCTTTAGCCAGCTGATGCCGACAGCCGACAGGACTTCGCAAGTGGCCGAGCAGTGGCCGGCGGATAG
CCTCAACTACGAGCGGCACTACAACCTGTCTTGGACCTGGCCCTGCTGGTCTTCTCTGCTGGCTGAGCTTCACTATCTTTTTATCACCCGACCCCTGTACGTGAGC
ATGCGCAACGAGCGAGCCATGAATTTGGCAGCAGTGGGCGGAGGTCAACCTCTCTCTGCGCTGTGCGAGAAGCCGGTAGCCAGCGACGCCAACGGAAGCCATT
GTCAGTCCCTCGACAGATTTGTGCCACACGACACATCTCACCAGCAGCACCACCATTCCAGTACTATGATTACGGCCACTGTGGCAGCAGCGGCGGATGAT
CAGCGGAGGAGGACCTGCGAAGGAAGGAGCATCTATACTGATGCGGAGCGGACTCGTGAAGTCCCTGCGCCATCAGCAGATCAACGAGGAGGTGGAACCCCTAAGTGA
ACTGGAGCTGGAATGGCAGTGTGTCAGCCGGGTGCACAGGATGCGCCAAAGATGACGCTGACGAGCAGCAAGGCTACGTGAGCGAATGAATCCCGCGCAAGTCAATTGT
CGCAGCCCAAGTGTGCTATCACGGAAGCGGAGTCTGCGAGATCCAACGAAACCAAGATCTGAGAGACAAAGCGGTGGTCAAGATGCTGTTCTGCTAGTGTGAGGAT
CTTCACTGCTGAGACCGCTGTATGTGATCAACAGATGACCATGCTTGGCGGAGCGGTGACGAGTACGTGGGCTATACCTCATCACTTCTCCAGCTGCTGGCC
TACTCATCCAGTGTGCAATCCGATCAGCTACTGCTTATGAACGCCAGCTTCCGGCGCGCTTCTGTTGACACCTTTAAGGGATGCGGGTGTGCGAGCGCTGTGCGCTC
CCTGCTGCTTCTGGCGGCGAGCTCCAAGAACGAGACGAATCTTTCGGTGGCCGGTAACTCCATTGCGCTGGCCAACTCGGTCTATGTCACGCCACAGATCTCGAGAGTCC
GCGACTCTGA
(SEQ ID NO: 62)

Start ATG: 1

MLPRLCADACRQCFAKIARRDTHRGTRTPYGCADTQSRPKPNFLLREVDEVCCCTAASAPRLLVLFDRDHKRASFFGLTIDAFYHYLRQALPLAKEAAIHLNASNEISAVGDG
VTITGTPGOLLNYSLELDGLDLNLDLMTLTPSSSTLAPAVTVTPGNRSVVRVSDVPVWVPCYSAILLCVAVGNLLVVLTVQNRMRITITNVFLNLIAISDILL
GVFCMPVTLVGLLLRHFI FGLLCKLIQFAQVPIPVTEPHNRLSNGHEPARPRADSPADRVGQHSADKEDDTSLLRMYRYAGNGVALHARLPGLQYLSIHGLGSAASVA
VSSVTLVATSCERYAICHFLRSRTWQTHANKI IAIWGLSLVCMPTIAAFSQLMPTSRPLRKCREQWPADSLNYERAYNLFDLALLVPLALLSFTYLFITRTLYVS
MNRERAMNFGSSPEVTTSSSAVAEAGSQRRANGSHCQSLDTIVPHQHNPHQHHSQYIYDYGHCGRRLISGGGPCEGRRLHYLCMRSAVSKSLRHQQINGGGTSLG
TGAGNGECCSRVHRMQMQLQQQYVSDNESRRKSLSQPSLRITAEGLRRSNETKSLESKRVRVRLVFLVLEFFICWTPLYVINTMTMLLGPVYVEYVGYTSISFLQLLA
YSSSCCNPTICYFMNASFRFAFVDTFKGMVRCERLCAPCCFWRRRSNETNLSVAGNSIALANSVMSSHTILESPL*

(SEQ ID NO: 63)

Name: GASTRIN/CHOLECYSTOKININ RECEPTOR like
Classification: G_{protein linked} receptor

Celera Sequence No. : 142000013384801

AGCTGTAAAGTACTACTACTACATATGTGATCAATTCAGTTTCGCTTACGAATGCAACGTTCCATTAAAGAAGCCACCAACTTTAAGATACACGCATATAACAACTCCTGAG
AATTGTGAAAACGATGATCTACTTTGATGATCAACTTTCATTGCTGCTGATTGAATAATTGAGTAAATGAGTAATTGGTATGCTGATAAACAGGAAAACCAAATGTCAACAGTT
GATAAGGAACATTAAAAACAAATTAAGAGGACGGTAAGTCTTCACTCGGACCTTTTCAGCTGAGCAGCCGTAAGTGTGTAAGGGGATTGCCCCACACCCAGACCGCGCC
ACTTCCACTTCCACTTTAAATAATCATATGCGCCCTTCTTTTGGCCAGGCTCGGCTGGCTTTGGCCCTTAATTGTGCACTCAGTTGCTGTGCTGTGTACTCGACA
AGGCGACGAACCGAGAGCCTTGACAGTCCATATGACAGGCGGATGGACTCGAACACTCGGACCCGAGCCCGGACCCGAAACCCGAGCCAGTCCACAGCCAGGACATAAGCG
GCTTTGCTGCCACTCACTGTGCTGCTTTCGGAGCTGCTCCTAGCTTGCAATTAATAATACCACTTTGGCCCTATAAATTGTGAACGTAGGAAGGCGGAAAAA
AATATTCAGCACTTTGGTGTATGGAGTGGGGTCCACTTAATTCAGGTACTCACCAAAAGGATTCCAACTATAAGACATACAGCGAGAGTGCTTAAGTTTACTCTGT
TTATTTGACGCAACCTTAGGAAGTTCAATTGAACACCAATAGACTAGCGTCTGAGTTATTTACGAGTACGATCATATACAGTAGGTAAAAAATCCATATTCGCTT
TACTTAAACAAAAAAGCAACACGAGATCGCTTATCATATATTTTACAGCGATCTCGTTGATCCTGAATGGGTCTTGGAGGCGAATCCCTCTACAGCTTAGAGCT
GAGGACTGTTGACGCGCGCGCGGCGCACTATTGGTGGTACACCCGCTTGAAGAACTCGTGGCGTGAAGGACTCCGCGAGGTGCCATACCCATGGCTAGGCGGGATTGAG
ACTAATGTTGGTGTGCGCATGCGATAGGCGCGCGGCGCTGCGCGCGCGCTGGCTGGCGGAGAGTCTCCACAGCAGCAGCAGCAGCGCTCCGCTGGCACTTGTCCACGA
CGCCAGGCGGAGCTTTAAGGTGTGACAAAGGCGCGCGGAGCTGGCTGCTCATGAAGCAGTAGGTGATCGGATTCGAGCAGCTGGATGAGTAGGCCAGCAGCTGGAGGA
AACTGATGGCGGTGTAGTCGACATCTGTAACACCCGCTCGATCAGCATGACCATCGTGTGATCAGTACAGCGGAGTCCAGCAGATGAAAACTCCAGCAGCAGGAC
GAACAGCATCTTGACACACGCTTCTTGTCTCAGGCTTGGCCCTGTTGGAGCTGGATAGTGGATTGTGGAGAAATGGGAATAGGGAATAGGATTGGCAGAG
CGGTATCGGACAGAACTCTAAGTGTGCGAAGTGGCGATCTGCGGAGCGGATGCTGGCGAGGAGTCTTGGCCAGCGTCACTGCTGGTGGTGGTGTGATCAGAT
TGGAGCGGTGCTGCTGTTGCAAGTGGTCTGTCATATTTGGTGTGATCCTCCGCTCGGATCCCTCTGAATTTCCATTATTATTGTTACTATTTCTATGTAAAAA
AAAAACTGTGAACAAAGCAATATAATCAACAAATTAACGTTTATAGCAGCAACAAATAAACAGTTGGCATACTAATCTAACTAATACATAGATGAAATAAT
AATTTTCAATTTGGTTTAAAGTCTCAAAATAATATAGCATATTTTACAGCGATCTCGTTGATCCTGAATGGGTCTTGGAGGCGAATCCCTCTACAGCTTAGAGCT
TGCCATATTACATGTGATGAACATTAATAACTTATGAGTTATGATATTTAGATGCAAAATAACTTGAATGCATTATAAATGCTCTTAAAGTTGTTGATATTCATG
GCTTTTAAAGGATGAAAAAGGATTAGTTTCGTGAGTTACCAACATATAGATGCGGTTGGCGGTGAGGAGGAGTACGATTTACTACTGCTGCTGGTGGCGGATTCGGT
GCGCTTCCGCGCGCGGTGTAGCGGAAACAGGACGAGTTCGTGAGGATGCTGCGCTGCGCTTGGCCATCGTACAGGTACGCGTACGATACAGGTACGCGTACGCGCAGC
AGAGGACGAGAGCGGACGAGCAGCAGGAGTCCAGCAGGATGTTGTAGAAGAGCTCGTATCCTGCTGGCGGCAAACTCAGGCAGCTTGAGTACGCTGTAACGTA

FIGURE SHEET 32

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Exon: 4080..4056
Exon: 2717..2517
Exon: 2454..2169
Exon: 1736..1001
Start ATG: 4080 (Reverse strand: CAT)

Transcript No. : CT21314
ATGCACCTCCAAACGGAAACAGGTGGAGCCGCCCTCCGTGGCCGTTTCGCTCTGGACCTTGGTGGCCATATCCTGTGAGCGCTACTACGCGATATGCCATCCACTGCGCTCGCGAT
CTCTGGCAGACAACTCAGTCAGGCTTACAAAGATCATCGGCTTCATCTGGCTGGCCGCGATCTCTGCATCAGCGCCATAGCGGCTTTTACTCAATTGATACCCACCAGTCGACC
CGGCTAGTCAAGTGGCCGTGAGTTTGGCCCGACACGGGATACGAGCTCTTCTACAACATCCTGCTGGACTTCTGCTGCTCGTCTCTGCCGCTTCTCGTCTCTCGGTGGCC
TACATCTCTATCAGCGGTACCCCTGAGCTAGGCTATGGCCAAAGGACAGCGGACGCATCTCTGCAGCAATCGCTGCCCTGTTTCCGCTACAAACGGCCCGGGAAGCCGACCAATC
CGGGCACCAGCAGCAGTAGTAACCTGCATCTTGGTCTTGACCGCCACCGGACTCTATAATGTTGGCGGAGGTAACCAATATGGCAACGACCCTTGACAACGAGACCAA
CGGCTCAACTGTGATACCAACCACCACGAGCAGCCAGGTGACGCTGGCCAGAGACCTCTCGCCGACGATTTCGCGTCCAGATGCGGCACCTCGCAGGTAGGATTCTGTGTC
GATAACCGGCTGCGCAATCCTATTCTCCTATTCCATTCTCCACAATCCAAATCCACTATCTCAGGTCCAAACGAGGCCAAGACCTTGAGAGCAGAGAGCGGTGGTCAAGA
TGCTGTCTGCTCCTGGTGGCTGAGTTTTCATCTGCTGGACTCGGCTGTACGTGATCAACAGATGGTTCATGTCGCGACCGGTGTGTTACAGGATATGTCGACTACACGGC
CATCAGTTTCTCTCAGCTGTGGCTACTCATCCAGCTGCTGCAATCCGATCCTGCTTCATGAACGCCAGCTTCCTCGGCGGCGCTTGTTCGACACCACTTCAAGGGTCTG
CCCTGGGCGCTGGAGCAGGTGCCAGCGGAGGCGTGGTGGTGTCTGCTGGTGGAGGACTCTCCGCCAGCCAGGCGGGCGCAGGCCCGGCCCTATGCGAGTGCCAAACACCA
ACATTAGTCTCAATCCCGGCTAGCCATGGGTATGGGCACCTGGCGGAGTGCCTACGCCACAGAGTTCTCAATGCGGTGGTGACCAATAAGTGC CGCCGCCGCTCAA
CAGTCTCAGTCTAA
(SEQ ID NO: 65)

Start ATG: 1 (Reverse strand: CAT)

MNSHNTGGAASVAVSSWTLVAISCERYYAICHPLRSRWQTISHAYKIIGFIWLGGLCMPTIAVFSQLIPTSRPGYCKREFWPDQGYELFYNIILDFLLLVLP LLVLCVA
YILITRTLTVYGMADKSGRILQOQLPVSATTAGGSAPNPGTSSSSNCILVLTATAVYNGWRRINQYGNHDLNETNGSNCDDHHHDDHGDAGQDLAQHSRPRCGTSQVGFVS
DNRSANPIPYSHFTIHNPLRSNEAKLTLESKKRVVKMLFVLVLEFFICWTPLYVINTVMVLIGPVVYEYVDYTAISFLQLLAYSSSSCCNPITYCFMNASFRRAFDVTFKGL
PWRRGAGASGGVGGAAGGLSASQAGAGPAYASANTNISLNPGLAMGMGTWRSRSRHEFLNAVVTNAAAAVNSPQL*

Name: GASTRIN/CHOLECYSTOKININ TYPE B Receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384546

[illegible]

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Exon: 1001..1564
Exon: 4101..4298
Exon: 4447..4618
Exon: 5695..5831
Exon: 5905..6087
Exon: 6381..6534
Exon: 7983..8249
Start ATG: 1149

Transcript No. : CT21432

Transcript No.: C124352
GGTTTGGCAACAGATTGGCAATATAGAGAGTCATCAAACGAGAGCAAAATGTCATACAGCATGTGCTCCTTATTCCTGGCGCTAGTGGCAATTGGAGTTGGAACTAGG
AGCCACCACTTAGACTCAACACCGCCGTGACCAATTCAACATGACGCTACTCGAGAGGCTTTCAGGCCATGTGCGCCACCACAGCAATAGGAACTACTGGGGGCGAGCATCAGCAGTT
TTGGTGGCGGAGCAAAATGAGCCTCTGGCGAGCAAAAATCCCGTCTCTGAGAGGATCAGCCTCATATGCCAGATATTGAAATTCATTCCGACGAGGCTCATGGAGCAGGGACT
GGGCACTGCGGTGGGCACTGGGAGCAGCATCGCCGTATCCGTTGAAGAGCTGGTCGCGCCGACAGGCCGAGGCATCCAGCGCAGCGAAGGATCCACCGACGACGCCAGCGG

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AGTAGCCATTGGCATTAGTCTTCGTCGAAGTGTTCATTATTGGTTTCATCTACTGGCCGCCATCTGGGCAACATGCTGGTATTGTGCGGTATCGGGACCGGAAAT
TGGCATCATTTACCACTACTTTGTGGTCTCTCTGGCCGTCGCCGACATGCTGGTGGCCCTCTGTGCGATGACATTTAATGCTTCGTCATGATCTCGGGCAAGTGGATGT
TGGTTCGCTGATGTGGACATGTGGAACAGCTTCGACGCTACTTCTCCACCGCCAGCATCATGCACCTCTGTGTCATATCGGTGACAGATACTACGCCATTGTGCAGCCA
CTGGACTATCCACTAATCATGACACAGCGACGCGTGTTCATCATGCTATTGATGGTGTGGCTATCGCCGCGCTCTCTCGTTCCGCCATCTGCTCGGGATGGTACACAA
CGACCGAGAATAACAATATCTCAATCGAATCCGCATATATGCGAGTTCAAAGTGAACAGGCATACGCCATAGTCAGCTCGTCGATGAGCTTCTGGATTCCCGGCATCGT
AATGCTGTGCTGATGACTACCGCATTTTACCAGGAGGCCGACCGACAGGAGCGTCTGGTGTACAGATCCAAGGTGGCCGCTCTGCTGCTGGAGAAGCATCTGCAAAATTAGCCAA
ATTCCTCAAGCCCGCGCGAGCATTCAGGTGGAGCAGTCGACCATCTCGACGATGCGCGGTGAGCGGAAGGCCGCCCGACCCCTGGGCATCATCATGAGCGCTTCTCTATCT
GCTGGCTGCGCTTCTCTCTGCTGATCATGCTATCTCTGCTGTGCGATAGTTGCATCCTCGCGCTGCTCTGTTGGCATCCTGTTTTGGATCGGCTACTTCAACTCGGCCCT
GAACCCCATTTATGTCATCTTCAACCGCAGCTTCAAGGCGCGCTTCAAGAAGACCCCTCAAGAGTCTGTTTTCCCTACGCTTCTACTTCTGTCGACGTGGCAGGGGCGGA
GACGATGACCGGAATCTGGAGTTCGGCGGTCCCGGCCGACGCGGGAACCAATGGAGGCCAAGCGACCGGATCGGATCCCGGAGATGGCCAACTCGCTCAACTCCACGCGCT
CGTCGGAGATACACATGAGCGTGTGCTGCCGCCAGTATGCGCTCAATGTCACACCCACCGGAGGCCAGATGCAGCAGCTGCATCCCTGTACACCAACTAA
(SEQ ID NO: 68)

Start ATG: 149

MTLLQRLQAMSAATTTTRILEGSISSFGGTTNEPLASKIPVLEESASHARYLKFIADGLIDEGLSVAVSGSSIIVSVEDVAVQAQDIQASEGSTDDADGSSHLALVFVKCF
IIGFTIILAILGNMLVIVSVMRHRKLRIITNYFVVS LAVADMLVALCMTFNASVMISGKWMFGVMDMWSFDVYFSTASIMHLCCISVDRYYAIVQPLDYPLMTQRRV
FIMLLVMWLSFALLSFLPICSGWYTTTENYKYLKSNPHICEFKVNKAYIVSSSMSFWIPGIVMSMYRIYQEAQRERLVYRSKVAALLLEKHLQISQIPKPRPSIQVEQ
STISTMRERKKAARTLGIIMSFLICHLPLFLWYIVSSLCDSCTPRLLVLGILFWIGYFNSALNP I IYAYFNDRFRAAFKTLKSLFPYAFYFCRRGRDRDDRDLEFGGPGS
RRGTNGAQRGTSGSAEMANCVNSTASSEIHMSVMRQYAVNVPTTDAQMQLHLPLTN*
(SEQ ID NO: 69)

Name: 5-HT1A/Ocr-like receptor
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013384515

CTGGACGCCGCGACAGAAAGCAGATCACCTGCCGTACAGTGCCAAAGATCATCGAACGCATTTACAATGAGGAGATCGGTGATGGCGGTGGCCACAGTGACGAGGAGATAAAC
ATGCTGGAGTTCACTCAGTATCTGGAGCAGTACCTGTGGCCACACTACCAACGTGAAACCGCCACCCATGCCACCTTATGTCCATCGTTATAATGGCCAAAGGAGAGTTCC
GGGAGCGCGTCAAGATTGGACCGTGTGTTGAAAGCTCCCGGATCAGTATCCAGCCTTCTTTCGCCACGCTGCTAGAGAGCTGTTTGGCGAGCAAAAGGCAAGGAGGCTAG
TAGCAGCGCTAAGGGAGCGGACGCGCTGTTGATGTTCAATTAACCACTGCTTTAACAGCATGGAGATAGAGCTGTGCAGAGAACAGGCCAAACGACTAGTCTCTCTCCATG
TGGCATTGCTGCGACCTCGTAAGTTCCTTTCGCACTCTTCACATCTATCAAAAGTATCAAAATGATTTGCTTGCAGGCGCTCGCGAACAAGAGCTTCGAGAAGTTCC
TGAGTGGCGAAAGTCTGGAACGCTTGTCTCAAGAAGGAGAAAGACAGCAAAACCGGAGGCTCTGCGGAGAGACACTTTATGCAAGACCTTATCATGACTTCTGCACTA
CTCGAAGCATTTCCCGCCGAGGCGAGGTCGCGCAACGTGGTTCACTACTGTGAGCGCTTCCCTAGAGTTTATATGCACTTGGAGGCACTTTTGGCGACTCGACGCTTCT
TCAACACGCTTTTAGACGACTGTCTATGATGTTGCGGGCTCTTCTGTCACCTTGGTTGCTGCGGAGGAGGAAACTGTTTGGTCAAGTAGGTGTGAGACTTCTCATTTA
ATAACTATCTCGTACAGCATAAATAGTTAAATTAAGTACAATCTCAATGTTGGACATTTGCTAACTACGAAACACTTATATTAGTGTGGACGCTGGCTCTCAGTAAT
CGTTGCGCTTATATAATGAGCTCATCGCCAGCTCGTTTGGCGAGCGTGGCTCATTTAGCAGCAGGCTCTTTTACGAGAAATGTCACATGTCGCTGGCGCAGCAGCA
TATGTATAGCAACAGCGCGCTCTGCAAGGCGTTAACTACGATGTTGGCGTAAAGCAGGCTTCCATCTGCAACAGGACATTATAGGAATAGCCAGGCTATAGACATTACT
AACAGCATACGCGAGAACATGATAAACTGCAAGGAAGATAATGTTTTATATACTTATATGTAATGCCTAGATGCCGATCTACTTACTTGGCTTCAATTTGTGCGCAATG
CGCGCTATACGCTCTCGGCTTAACTCACTTCTGTTGTCACATAGAAGAAATGTTTACCAAAATGGTACAGCAATTGGAGCAAGAATATGCAATATGCTAGCCAGC
CGATCGTCTTCTGCTCGCCACCATGTGCTCTGTTGTAGGATTGCGCGTGCAGGAAGAAATGTGCGAAGACAGCAGTGCAGCCATCGTTGCTGTGCATCCCCAAGCATA
CGCAGAAATATAGCAGTACTTCTGCGCTCCGTGACTCGCAGGAAGACGTTCTCGAAGCAAGGCTTCCAAATGTAAACCGAACTGTTACGCAAGAAAGCTGCC
AAGAGCTGAAGCAGGATAATGTCCGCCACAATGAAGCTCACATGGCTGCTGAGCTCGGTGAAATTTCTGACACAGGCTGCTGCTGCTGGCTTACCATAAGGCACATGGCTA
TCGTTGTCAATGTTTACCAACAGATCTCTGCTGAATGGATTGAAATTAACGCATGAAGCAAGTCAAAATCAAACTATTGCTCACCTGAGAGTAGGAAGTATAAAATAGA
TGATGGCGATGACGAGCAGGATAACAAGCGAGATGCCATGGAAATTTGGGTTAAGATTTTGGCGAGCAGAAAGTTGAGTCACTCCACTTGATCTCCTTGGCGGCAAGCA
AATGTTGGCGAATAGCACATTTTCTCTCAGTACTGGAAGTGGCCTGCGAATGGATTATGATCATCCATGCAATTTATGTTGGACTTACTACTCTTACCTTGTCAATG
CAATACATCCCTTATCGTATGATGTTATAGCGGTTCCAGACTGCGGCAATATCTCTCATATCCGATTGATTTCCATGACGCTCTCTGCTTCTGTTTTCGGCATTGG
TGTAATGCTTACGCTCCCATCATCCAGTAACACTAGCTTATCAGAGCTCTAGAAATATAAAATATCTATAAGTATCTAAAAGTTCTATGCAAAAGACTTCAAACTCAC
GCCAGCATAGTGTATATTGGGCACATTTGGAACATTTGGAAGTCCGATGAGCAATTTGAACCTTAACGGGCTGTTTGTCTCTCTCATAGCTGGTGAACATC
GGCTGGAAGTAGTCTGCAATGGAACGAGCAGATAAATAGAGAACACAGACCGTGGAAATCATAAAGGCAGTTCAAGTTAAGCTGCTTTTGTGCGGCTGGTGGATCAGA
GGGAGCGCAGATTTCTTCTCTGCGAATAAATATTACGCTTCTGGGCAAGAACTTGAGCTGTTCCATGCGCTTCTCATGCTGTTTGTCAATTTGTGCAGATT
GGTTCTTCTTCAATTTCCGATACATTTTTCCTCAGTACGCTTCCAGCAATTAATACCAGGCTCGCATTTATGGGGAACAGAACATCTTATATTACCTTACTA
GTTGATTTGTAGTATTGGGATTACTCAATGACTACAATAAAATCGGATGAGATCAGTACAATGCAATGATTCAACTATCAGTAATAGTTACATTTTATGTTGTCACCTAT
ATGGCCGATCCATGGTCATAAATGTTTATCCAGTTTATGTTGCGAGCTAGAATATCGATATAGATGCTAGAGTACGCTTTTCTATTTCACAAATGTTGCGATTGTTT
GTCAAACTTACCCAAAGGCAATATATCACTATATTCACCATTAGCTCCACCAACATTAATACCAGGCTCGCATTTATGGGGAACAGAACATCTTATATTACCTTACTA
ATGGATTTTTTGCTCATTTTACTTCCAAGAGATAATATCCCGATTGGAATCCCTCATCAAGTGGCTACTTTATAGTTTAAACAGAAACATGCCCCACAGCATAAATCA
TATTTCTCATCAGGATTGGGGCCACCTCTGTTTGTGTTTAAAGAACTTCTTGGGGAGTGCGAAGTGTGGAGTTGGGGATATTAGCTGTTGGCTTCTGTTCCATGATCCAC
TGATAGCCAAACAATGGGCCATTTAATGTTGGGCATACATATGTTGATGCCAGAGGGAATCCATAAAGTTGAATGGCCAAACCACTTCTGTTGGTGTGTTATGGCAAC
TTTTTCGAGCGACTAGGAGTTTTCGCTGCGCAGCTGTTGAACTTACCGCTTTCATTGACCTGCTGACACTCATGGTCCAGCTGTATCTCGAACCTTCGCAACATTTGTT
GACCAGCACTAGTTGGGATCGGAAGATAGGCTGTTGATGATCGGCGCTAGTGACATGACTGTT
(SEQ ID NO: 70)

Exon: 2651..2621
Exon: 2477..2353
Exon: 2289..2119
Exon: 2061..1880
Exon: 1824..1321
Exon: 1259..1001
Start ATG: 2651 (Reverse strand: CAT)

Transcript No. : CT21585

ATGGAACAGCTCAAGTTCTTTGCCAGAGCACTACTTCCAGCCGATGTTCAACAGCTATGGAGGAGAGCAAAACAGGCCGTTAAGTTCAAATTCGTATCGGGATTCCAA
ATTGTGGATCCATGCAGATGTGGCCAAATATACCACTATGCTGGCAGCTCTGATAAGCTAGTGTACTGGATGATGGGAGGCTAAGGCATTACACCAATGCCGAAAACGAAGC
AGAGGAGCGTCATGGAATACAATCGGATTATGAGGAGGATATTGCCGCGAGTCTGGAACCGCTATACCATGACTACGATAAGGGATTGTATGCAATTGACAAGGCCACTTCC
AGTACTGGAGAGGAAATGTGCTATTGCGCAACATTTGCTTGGCCACAGGAGATCAAGTGGAGTGACTCAAACTTTCTGCTCCGCAAAATCTTAAACCCAAATTTTCCATG
GCATCTCGCTGTTATCTGCTGCTCGCCATCTATTATTACTTCTACTCTACGAGAGATCTGGTTGGTAAACATTGTGACAACGATAGCCATGTGCTTATGTT

FIGURE SHEET 35

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Exon: 1693..1490
Exon: 1189..1001
Start ATG: 4299 (Reverse strand: CAT)

Transcript No. : CT21642

ATGACTGTGCCGACATTTGTGCGCATTTTCGGAAATGCCAAGGCCCGCGCCCATCACATCAACCGCCTGGAATTTTCCACCCCTCTGCGCACAACCCACCGATCGTTT
GGCTTTTCAGGCAATTTGCTATTTCAATTTGGTGTTCGACTTGGCAGCGTTTTTGCATTGTCCACCGAAATGGTTTGGCCTGATTTCCAAATGCGTCGAGTGCCCAACAGAAATC
AACACCATTTGCAATTTCTTTTCGATATTTCTGACGGTATTTTGAAGAGCGCATGAGATGCACGACGAAACATATCTAACGGCACTGGCCATTACGGATATTGCCTACCTG
ACGTGCCAGTTAATCTGTCTACTCCAGCACTATGACTATCCCAAGTATCATTTCAAGCTCTACTGGCAGCTCTATGGCTATTTTGTCTGGCTGTGCGACAGTTTCGGTTACA
TCTCAATTTACATAGCCGTGTGCTTACCATCGAGAGATTCAATTGCGATACGCTATCCGCTCAAGAGGCAACATTTCTGACGGAATCGCTGGCCAAAAGGTGATAGCAGC
GGTGGCCATCTTCTGTGTGTCTGACACATATCGACGGCATTCGAGCATACAATTACGATTGGTACTAGGCAGATTGATGACGCTACAGCCGTGCAACCAACCGGTGGCC
AACATCTCGCCCATGCGCGCGCCACCCGTGGCAGTCACGCCCCCACTGGCCACGCCCCCGCTGCCACGCGCGGACCATTTGGCAATCGCCGAGTCCGCCATGGAGTCCA
CGACGTCCGGCAGCTCTAATCAGCTCGTGCAGTGGGGCAGTGGCAGTGGTGGAGAGCCAGAGAACATTCACGACATCGCCGCTCACTGGCAGTCACTGGATTGTGAC
GCTGCCCACTGAGAAAAACGCTGGAGGAGCAGGATCAGAAGTGGCAGACGCGGCCAGAGGTGAGGGTTCACAGAGAGTTTCTGCAATTTGTGGCTCGCAAGCGCAGC
GCTGAGAAATCACAACTAACAAATACGGATGCTTCGCATTTAACGTAACGGAATCTGTCAAATGTGACCTATTACAATCATGGCTTTTCGAACTGGGCTATGATGAGC
TATACAGCTATCTGTGGAACCTTGTACCCTGCTGGTCTCGTGGTATTCCTCCCTACTTCTATTGGCCACCTTCAACTCCATTCTATTGGTTTCATCGGTCACAGAA
TCTGCTGTGACCTGACCAATGCCAGCAGCATAGGCGCACAAAATACCGAAAGAGCGTTTCGAGAACTGATAACGGGATATCGCTATAGACGTCGACATGCGGCTAACAAAT
ACGAGTCTCTATGTGCGCATACAACGACTACACTGACCCAAATAAATGGCGATCATTATGGTAGTAATTATGCGGAGCCGGAAGTCGTGCAATCGTAATACGGGTCCGC
TGATAGCGTGA
(SEQ ID NO: 74)

Start ATG: 1 (Reverse strand: CAT)

MTVPTFVRISENAQGPAPITSTAWNFPPLRTPPIVWLSGNLLFQLVLHLAFLHCPKWFGLISKVECPTESTPFAILFDILDGIFRKRMRCTTNIYLTALAITDIAYL
TQQLLSLQHYDYPKYHFKLYWLYGYFVWLCDSEFYISYIYAVCFTHLFIATRYPLKRTQCTESLAKKVIAAVAFCLLSTLSTAFEHTITGTQIDDAYQPCNQTV
NISPMPPPPVAVTPPLATPLPTPATWQSPDSAMESTTSGSSNQLVDWGSAGGDFENIPRHRHWQSSGFVTLPLRLKTLLEQDQKVADAAQSRGVTESSLQLWRRKRS
AENHNINNTDAFAFNVEYQCNVYNYHGLSELGYDELYSLWNLFLLLVFVFFLLLLATFNSILILLVHRSKNLRGLDNLASSIRRTKYRKTRELTGYRYYRRRHARN
TSLYVPHHTTTTLTQINGDHYGSNYGGAGSRNRNRTGLRIA*
(SEQ ID NO: 75)

Name: TRH receptor like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013384839

ATAAATGGCCGAATTTGCAAGTCAGTTCCACTTGCTCTTTAAAAAGCGAATCTCACCAGAAATTAAGCAAATTTATCACCGACCAATGTGAATTCCTACTAATAATTTTTC
AGCGGTGTGATGGCTTTATGACATCGCCAGAAATGATTGAGCTGTAGCAGCGAAATTTAATCAAGACCTAACCCACGAACTGGACGTTCTCACATAGAAAAATGAAA
ACGAAAGCTGAAAGACAGATGGATCGTTGATGTGAGCTCTTTTACACTTGGCTTACACAGCAGTACGCGCGTGGTAAATTAATATACAAACAAATTACACGGCCCGGC
AAAACAAATTTAAGCCAGTTAATGCCGTGCGACAGTGGCTGCAAGCCACGCAATTCGCGCGGAAAAATGGCCACAATGGCCACAATGGCAATGAAATATGACAGGGCGCGGTTT
TTGCATACTCTACTGTCAAATGAAGCGTAGCAAAAGGAAAAATGCTTACACATACTAATTTGAAAAAGAAATGCAAAACCAATGCAATGAAATATGACAGGGCGCGGTTT
GCGAAGCTGTCTAATTTATTATTGACATTGGCAATGACAATGAAACCGAGAGTCCACAGCAGAGTAATGGATTTTCTTCAACTGCTTAACTTCTTTTCATTGAAAT
CTTATTGGACATATAAACCATACGTAGAAATGATAAGAAATAAAAACAAGTATGTTGACACAATTTGATAAGGAATCTTAGTACTTTCTACCTAAAAAGGTGATAT
ATATTTGAATTCATGTTGTGATGATAGCTACTCGAACCAGTGTGCGTTGGCTACAGTCTGCTGATGCTCAGAGTGTGTAAGTTAAGACTTTTGCTTTCTTCCCCGACT
TGCTTTTGTCTTTTGAAGAACCAAGCAATGGATAAACAGAGTTCAAGAGGTTGTGAGCCCGCCCTTTGCGCAGATGAGGCAAGAAAGAAAGTAATATCAAAA
TTAACAGAGGCTGACAGCTGACAGGGCAGAGGAATGGGCGCAATGGAATCTATCTAGCTTGAATTTGGTCTCGACCCGCTCGGGCAGAAATGTGACAAATTTGAGGG
GAAAAGCTCGTATGAGCGGGACTTACCACCTCCATTGTGGCAGTGGAGCAGTGTGAGAGTGGTCCCTGGCCGCTGTTGGATGACAGATTGGGCAGTCCGACGCGCTC
ATCACGCCGTACTTGATGGCTTGAAGTGAAGAACGAGGGCAGTAGCCGGAACCGCGCGACTCAGGGAACTGAATGCTCTCGCGGCTCCACGTAACAGGCGCTCCT
TGCCCATGGCCCGCAGTTGCTGCCCTTGGCGATGAACATCACCAAGAGACCTGGCGGATGTGCTTACAGCTTACAGACCCGAGGCGCAGCAAGCTCCTTGTGCGCTCGGCCAC
GGCCAGGCGCATAGCATATCCACGCGAGATGGGATGGCTCTCCGATGGCCACGCCAATGTAGTGGCCCTACGGTAGTTGTCCCTGATGCGCGCGACTTGATGGCT
AGCACGGCGATGAAGAGCATCAGAAAGACAAATGTAGATCAGCGAGAAGAGCAGCTCCGAGAACTGCGTCTTGACAGCGGATATTAGTACGGTGTAACGCGCGCTATCCGG
TGATAGTCTCCGGAGTCCCTGCCAGCGTTGTGTAGGACGTGGGTAACAGAGTCCGAGTGTGCTGTGAGGCCACCCGTGGTCTGAGGAGCCACCTGGTCTAAGGAAGCCACTGCCATCACAGG
CACACTGGTGGTGTATACCCTCCGCTGGTGGTGAGCAGCCACTGCCCTCCGATCGCTACCTGGATGAGCAGCGCAAGAGGAGCAGCAATCCCTGGTATGGAGCCGCGCAGA
TACACCCCTCCATTTAGACTAATCAGAAATACGCAATTTGACAGCAGGGCGGCAAGACAGGGCGTAGGCCACGCCCACTCCGAAGCGGATGGCCCGCAACTGATCAACG
AGGGCTGTGCGGTGATGATGCCCCCAACTAGCGCAGGCGAAGACAGACCGAGAAGCAGCAATTCGCCGAGTGTGCTTACAGACCCGAGGCGCAGTGGCAGCAATTTAGATCA
GAGCAAAAGATCTCGAACCGTGCCATCATGAGCATGGTGAAGGTGGCCAGGACTAGCACCAGGCAACCCACGCGCTCTTGGTGTAGTCCCGCAAGGGCAAGTGGGTGGAC
ACTATACTGGCGGTGTGCTGTTGGAGCTGCTCCCGGAGTACCCGCTCAGTGGTGGCGGCTGCTCGCGCTTCGCCCCGTAACCTCCGACGGGGCGTGGCTGGTGTGGTGG
TGGACGTGGTGTGCTAGAACTGCTGGCGATAGTGTGGTACCAAGAAATTCCTGACATTTCGAGACGGGCGAGCACTGGAGACATTGCCACCCGGAATTTAGATCA
CAATTTGTTGGCAGATCGCTGGGTCTGAGGAGCTGCCGAGTAGTGGTACTAGTGGTGTACTTGTAGTACTTGTAGTCTGCTGGTGTAGTGGTGTGCTGGTGGCATT
CGGCTACTCGGCTTAAGCACCTGGTAGCGTTGCGATCCGATTCTCGCTGGTCAACAGGCGCACCCGCGCCAAATGTGTTGCCATTAACTGAGGCTGCGCGACGCGG
CGGTGGTGGCATCACTGCATGATCAGCAGCAGGATCAGCAGCAGTGGCTTACCGGCCATTGGATGTGGTGACCTTCGATGCCCTGGACCTCTGACAAATATAATCAAA
CGCGGCACATCTCTTGCACACGACACTAACCCGCACTCGCGATCGCGATTGTTCAAGTTCGTAATTTCTTCACTCTCGCAGGCCCTAGATCATGAATTTCTTCTCGA
TTCTCGCGCTAAACGCTCCGCTCGCTCGATCGACCAACTGCAACTGACTGGCGGCTCTGCAAGTTGGGCTTTGGACCGGCTTAGTCTGTTGAGACAGTTTGACGGCAGCC
AAGTCGAACAACTTGTGTTGAGTTCGATGTGCGTTTGTAAACCCGCTCAACACTTACAGGAACGCAACATGATGCTGGTGGGAGCAGTGGGTTGGGGCGCTTTATATG
ATTTTTGCAATTTTCTCTCGCGTTCGAGTCTTGAATTTGATACAACTTCGTTACTCGATCTTCTGGCTTTAAGGTTAGTATATAGCTTCAACA
ACTAATGTTCTAATAGCTCAATGTCTTGTGAAGGCTATTGATTTCTCTGAAAGCTATGTACTTAATTTCTATTAGACCGAGTAAGAACTCAATTTATCTTTAGC
TTCCGTGCTAATCCGCTGGAACATATACTTTAACCAACATTTACATACGTAAGTTTTTTTCCATTTGATAATTTAGATTTAGATTTAATCAATTAATCAAT
GGACACAAATTTTTCTCTCGCGTTCGAGTCTGAGCAGCTTCAAGCAGCTTACAGACGCTAGCTGAGATCCACACTCAAACCGACTGCGGGATGGAGAACACTTTTCAAT
CGTCTGCAATATCGGAAATTTGGTTTATGGGCTCTGATGCTCGAGTGGCTGGTAGGAAAGTCTATGGGCCATTATCTATTAGCCAAACCGCTTTTACGGTCTCTCGCGG
AGCCTCTCTCTCGAAACGAGCCAAACAAAGGCCAAATAGCGAAATCCGAGCAACAAAGGTGATTGTAATCAAAAGAGCGCTAATGGATATATGATGATCAAGAGCTGTGA
GGCGGAAAGGCAATCGCCAAATAGCGTTAGAAACGGCAATAGCAATAGCAACCGCTGATTCTTCTAGTCTTATGTTATGCGGCGCGCTTTTCACTCAACCGCAGTTAGA
ATTGATACCTGAATGAGTGGGCA
(SEQ ID NO: 76)

Exon: 2946..1001
Start ATG: 2752 (Reverse strand: CAT)

FIGURE SHEET 37

38/89

TCGATCCGAGGCGACGGCGACGTTTATAGCGCGAGAATCGAGAAGAAGAAATTCATGATCTAGGGCCTGCGAGGAC TGAAGAAATTTACGAACCTGAACAATCGCGATCGCGAGTGC GTGGCTTAGTGTGCGCTTTGCAAGAGATGTGCCGCGTTTGATATATTTGTAGAGGTTCCAGGGATCGAAAGTCAACCAATCCAATCGGCGGTAGACGCCACCTGCTGCTGATC TCGCTGCTGATCATGATCGATTTGATCGCCACCCAGCCGCGTGTGCGCGAGCCTACGCGTTATGGCAACAACATTTGCGCGCGGGTGTGCGCTGGTGACACCGAGGAATCGGATG CGAACCGCTACACAGTGCTTTAGCCGAGTAGCCGTAATGCCACCAAGCCACCACCTACTACCAGCACGACTACAAGTACTACAAGTAGCACCACTAGTACCACCTACTCCGGC AGCTCCTCAGACCCAGCGGATCTGCCAAACAATTTGGATCTCAAAATTTTCGGGTGGCAATGTCTCCAGTGTCTGCCGTCTCGAAATGTCTGACGCGGAATTTCTTTGGTATCCACCA CTATCGGCCAGCGGCTTAGCAGCAGCCACTCCACCCACCAACCCAGCCCGCTCGGAGTTCTACGGGGCAGAGCCGAGCAGCCGCGCACTGGACCGGTTGACTCCCG GGAGACGCTCCAAAGCAGCAGCCGCGATAGTGTAGTGTCCACCCACTTGGCCCTTCGCGGGATACGCAAGGAGCCGTGGGTGGTGGCGGTGCTAGTCTCTGGCCACCCCTCACCAT GCTCATGATGGCAGCGTTTCGAGATCTTTGTGCTCTTCAAGGCGCTGGCGGACGCTCCCGCTGCGGTAGGCACCTCTTTCGGGCCAGATGCTGCTTCTCGGTTCTGTTCCGCTGC GCTAGTTTGGGGCCCATCATCCGCGACACGCCCTGTTGATCAGTTTCGGGGCCATCCGCTTCGAGGTGGCGTGGCCCTACGCCCTTGCTCTTTCCGCCCTGCTGTGTCAAAT CGGTTATTTCTGATTAGTCTAAATGGAGGGGTGTATTCGCCGCGCTCATACACAGGAGTGTCTGCTCTCTTTTGGCGTGTCTATCCAGTAGCGATCGGAGGGCAGTGGCTGCT CACCCAACCCAGCGAGGTATACACACACAGTGTGCTGCTGATGGGCAGTGGCTTCCTTAGCACCCAGGTGGGCTCACAGACCAACTACTCGGCACCTGTTTATCCCCACGTC CACACACCGCTGGACGGGACTCCGGAGATCTACACCCGGATACAGTCGGCGGCTTAGCACCGTACTAATACCGCTCTGCAAGACGCGAGTTCTCGGAGCTGCTCTTCTCGCTGATCT ACATTGTCTTTCTGATCGCTTCTCATCGCGTGTGCTAGCCATCAAGTCGCGCGCATCAGGACATACCGTAGGCGACCTACATTGGGCTGGCCATCGGAGGCGCATCC CATTGCTGGCTGGATGGA TGTATGCGGCTTGGCCGTGGCCGAGCGGCACAAAGGACGCTTGCCTGGCGCTTTCGGTCTGGTAGGCACATCCGCGCAGCGGTCTTCTGTTGATGTTC ATGCCCAAGGCGAGCAACTGGCGGCCATGGGCAGGAGGCGCTGTACGTGGAGGACCCGAGGACAGTTTCAGTTTCCCTTGAGTCGCGCGGTTCCGGCTACCTGCCCTCGT TCTTCCACTCTCAAGGCCATCAAGTACGCGGTGATGAGCGGCTGCGGACTGCCCAACTCTGCAATCCAAACAGGCGCAGGACCTAGCTCAAAACACTGCTCCAGTGGCCAAACA TGGAGGTGGTAAGTCCC GCTCATACGAGCTTTTCCCTTCAAATTTGTACATTTTCTGCCCCGAGGCGGTGGACCCAACCTCAAGCTAGATACAGTTCCATTTCGGCCCCAT TCCTCTGCCCTTGTACAGCTGTGAGCCTCTGTTAATTTTGTAA

(SEQ ID NO: 77)

Start ATG: 195 (Reverse strand: CAT)

MAGRHHLLILLIMQLIATTPASRSLVNGNNIWRVRVLVTSQESDRNAYQVLPKSSRNATSTTTTTSTTSTSTSTSTTTPAAPQTORSAKQDLNLPFGNGVSSAARLE
MSTEFFVYVSLTASASSSTSTSTTPATPRRSSGAKARAAGTGRVTPSSSSSTPISVSTHLPFAGLRKEPWWVPVLVLATGLTMLMAAFEIFVLFKAWRTSPSRHHFLFG
QLLLGLGFACASLGGIIITAQPSLISCGAIRFGVQVLIAPVLAALVKCVFLISLNGGVLPAPYQQLLLFALLIQVAGQGWLLTPQPEVYTSVPMVSGSGLSTVTAQST
NYSALFYPTSTYTLDTGTEIYTRIAAVSTVLPLCKTQFSELLFSLIYVFLIVLFAVLAKSRGIRDNRYREATYIGLAIGGAIPILWLGWMLCGLAVAERHKDACVAFGLVA
TSATVFLVMFMFKGRGLAAMGKGLYVEDREEQFSSLSRAGSGYSFHFHFKPIKYGVMSGLPNSASNTGQGLSSKKHCSANNGGKRSRYELFPSPKVFYTSARGGRQL
KLDVVPFPHPSALVSCQPLLIL*
(SEQ ID NO: 78)

Name: Metabotropic glutamate receptor-like
Classification: G protein linked receptor

AATTTGGTATAACCTTCAGTCTCTATACTCAGTTATTCCATTAATCTCTAATCCCAACGACATTAAATACACATACAAATTTTTTATACAAAGTTTGCTTTATTGACTATTT
 TTAGAACACATTTACCATATCTTTGTAAGATTAAATACCTTAGCATAGATTACAGAATTTTCAGCTATTTCCTAGAAATCTTGCTGTCACAAACAAACAGATTTTACATGCTTTT
 GTTTTATAATCCCTTTTCTTTGGGAAAGATTCTCATATATTACCAATATAAATTTAAATAGTCTCTACTTTAGCTTACTCATATACTAAATTAATCTCAGGAGGCTGACT
 TTTCCGCGGACTTTTATGGTGGCTCTGACCAACATGGATGCATCTCGGCCTTTCTCGCAGAAAAGTATTTTATTTCACATTTTACGGGCGTCTTGCGTGGCTGTCTTACC
 TCTGGCTAAGATGCTTAATTTTATACAAATCCGCATTTTCGGCTTTTAAATGCATAAATCAGCTTAGTATTTAAATGGTTTATTAATCTACGGGGAACCAACATTTTATT
 TTAGTAAGTACCTAAGAAAAGATTGGATCTTTAAGATTCCGATATTATTATATCTTTTATTACTTAGTAAGAAGTAGGGGATTTACGTATATTTTATTAAAGGGTAT
 TTGGCAGTAAATATTTATTCAAATTTGGGGTTGTACAGTTTATATTAATTTAAATATGTTTATTAGCTGTCAAGGCAATAAGAAAATGTAATCCATTTACGCCCCCTTCAG
 TTAATATATTTGCTAACATACTCATGTGCGGGTGTACAGATATTTCAAATCTTTGAAATTTAAATCTGATCTCGGTTTTATTACGTAGACACATATTAATACCTGCTTA
 GTCCTTAAGTCTTCATTCATTAATTTGGATAAGAAAGGATTATAAATCATTTAAATGCACACTATAATATTGGCTTATTTAAAGGAAATTCCTACTTTAAATCGGGTTACAAAT
 CTGTCTGTGCACTGCGCCCGATCCTCTTCTCGCGGTGGTGCAAAATGCCTCCTGCTGCTCGCTCAAACATATAATCAAATCTCCGGCGGAGCTGTCACTTTGCCATTGGA
 ATTCGATGATTTCTGTGTCGCCGTGGTTCGGGTCTTTCCGCTGTCTGGTGGCAGGATGTGGTGGTGCATGAATTTGTTACCGGCATTCAGTGGGACACAGCTGGGAATATC
 TGGCGGATTCGTGGTCAACAGCGCTTGGAGCCACCCCTCTTTTACGCGCTTTTGGTGAARAACACGGGCTCCAGCTGGAGTGGAGGCTATCATCTGCTTATTCATAC
 AGGTAAGGCCCTTGAAGAAGCTCTTCCGGAAGTCTCACTTAGGAAGGCATAAAGTATGGGATTACCAGCGGAATTCGAGTAAACCAAGTGGCCCCAGAAGTAGGAAAATGAG
 TATTTCAGCTCGGGAGAGGTCCTTTCCGCGGGATTTCAGATGAATCAGGGCCACCTTGAGATCCAGTGGTGTCAGAAATGAAGATCGAATATCTTTAATTAATTAATACCT
 GAAATACCAAGTAGAGGAGGCAACATAGCAATGTATACCTATCAGCTTATCAGCTGGGATGAAAGAGGTGGGAATGTCAAAATAAATTTGAATCTTATTCCTGGCATATACCG
 TCAGTACCAGTCGAGTGACCTTCTGTGAGCGCGCTCTTCTCCTTGGACTTCGTTCTCGTGTGGGACCCACCGATCGCAGTTTCTTATAAACCAAGTAGTAGAACTCAC
 GATAAGAGCACGGCTGTGGCGAATCTTAGGAAAATGTGGTACAGATGAAGGTGGTGGGAAATGCTTCTTACGCTATCGGCCACATTTATGTGACAGGTAATGTATG
 CCATCTCCTCGTCCACAGTCTGGCATAGGAAGTACGGGACAGATGAGGACGGCTCAAGTTGACACGGCAGCTCGAGACGTTTGGCAATATGCAAGAGTCTGATATC
 GTGGCAGGAAAATCGGTGGCATACCGCTATATATCGATCCCGGACATGATAGGCAGAAAATCGACGAGGTGAAGGAGGTATGGATGTGCTCACCATCTAGGCTTTGCA
 CATAAATCCCGCAATCGCCAGCTGCAAAATCTGCATTTGTGTACACAGAAAAGGTTATCCAATCAGGAAGACATCTGCTGCCACCGCCAGATTCCAGTATATATATCTGCTG
 ACCGTTTGCATTTTGGAAAAGCGCAACACCGACTAGATCAGCAGGGTGTGGCAATAATCCGTAATATGCACACAAATCCGTAGACGACCAAGTGAACAAAGTACGCAAAAG
 AATTCGCGTGGCTATGCAATGTTGATAGTGGGCAAACTAGTGCATAAAGGAGCTCTCTCGCTGAATTTGTGAACATACCATATCATCTATGTTGTTAGTTCAGTTCCTGGT
 ATATAAGCTCTCGTCTGTTATACCAATTTGTGGCATTTGGCTCTGAAATCGGGTCTCATTTTCCCTTTCCAAACTGGAGACATCCATCATCAGGCCAGTAAACATATTT
 CCACCTAAGCGCTGCCAGTTCAAATTTAAATATTTTAAATAGGTTTAAACGGTTGTGCTCCCGCTCGTTTGTGTTTCGATATTTGGCGTTGCTGCAAAATACATTTTGT
 TATTACGTGAAAATTTTTCAGACACAGTTGTTTGGCCAACTTGATTAGTTACTAATTCGAAATACCTGTTGCTCTATTAAAGAAAACCACTTAAACCTTCACTAGTCTAGT
 CAATATTTTATTCACCAAGTAAATTTAAATGTACGTGAACAAATTTGGTGAAGTATTCAAATGGAATTAAGTTAAGTCTACAAGTTTATAAGATTTTAAATATACTA
 GATTGACGCTCTTAAATGCATAGTGTATATTTTAAATATATAGATTTACTTTTAAAGTGGAGCTCAATTTTATGCTGTCTTAACATCTTTGAATCTGATGTGACCCCAT
 TCGTAATGCCACTTCACATACAGATTCCTTTTGAACCTTTACTACAACTTTTACTGACTTTCTGAGAAATCCCTCTGAAAATGAAAACCTTCTGCAACACTTTGTAGAAAA
 CAAACCTTTAAAAATACAGCATGTGAATTAATACGTCTGTTTCTGCTTTGAAAAGTGAGTTACCAACAAATTTGGTTTGAAGGCGAAGGAATTTAACTATATATTTAA
 TTAATTTGCTCAGCTTTGACAGCTGCACAAACCAAGAAACAAAGATTTTTTATATTCTGACCTAACCTGCACACTTCTGTAAATTAATTAGGCTAAGGAAAAGCTGGCG
 AGCATTTATGCAAAATACAGGTGAAAATATTTGTCTTCTCGGAAGTCGGAATACTCCAGCTATACAGATCCCATGGTTTGTATTAGGAGATGAAAATTAAGCCGATTTG
 TTTGTGCATTCATGCACATCAAATATTGTTGACTGAGCCAAAAACACATGTG
 (SEQ ID NO: 79)

Exon: 2524..1679
Exon: 1510..1001
start ATG: 2524 (Reverse strand: CAT)

Transcript No. : CT22465

40/89

CTCGGGAATTTTCCCTTTTTCGGCTTTTTCCTCGTGCTGAGGAAATGACGAGAGCCGAGGCCAAGGAAATGAAAGAAATCTTTTCAGTTCAAAGAATACAAATCCATAA
GGCGAGAGAGATACATACATTCGCATATTTCCATCCAGGACGACATTTACATAAATACACGCCCTTTTCCTCCGCCGCCACATTTGCACACACTCTTCCTCGAGTCTCGTTCGTG
TCGGTGCTGCTATTTCTCTGCTGCTACTTTGTGGGATACACTTTGGAGAAAATATATATATGCTAGTCTCTCTCAAGAAATCCCTACACATATGCTCTAAACAGCAATCTCA
TTGATATTTACAGGTTTCGAAAGTTCCTTAAATACCTCAAGAAACATTGCTACTACATTTTACTTTCATGAGATACAAATACAAAGACTTGTATTGGATTGTCTCAGTGC
ATTGCTTGGTTCTGGGCGCCAAAGTTTGTGAGCAATTTTGTAGGAGGATTCACGGAATGGCATCTTCATCTTTGTAATTCACGGCAAGGCGCGGCTAGAGGCGCTCTTT
CGGCGAAGGACTCTAAATGCTTAAATGGCGCTGACACACTCACACACACTTACACACTCACACTACACATAGACTACCTATGAGATGGACGGCGGACGGCGGG
GGCATGACGAAGATGCCACAGGAAGGTCGGCCACCGCCAGGCTCATCACAAGCAGTTGGTAATGGTGCGAAGCGCGGAGTGGTCAATATGGCCAGTATGACCAGGGTGT
TCCCGATGACGGTGACCCAGTGAAGGTTGGCAAGGATGCAATCAGGATGATGCTCGGTCGAGGAGAGCAGCAGGATGTAGAGATGGCCCATTTCCGCGCGCGCCGACCT
GTTGTAGTAATTAATGTGGTCACGGGCGCGGTGGCTCGGGTTCGGTAGAGCCACTGCATCCGATACGTTTCGGTTCGGTGGGATACCACTCTACTCTCTCTACTCTCT
CCTCCACTGCTCCCATTTGCCCGCATCTCATATACGATGGATATGTTTTCGGCGAGACCACCCTCTCCCACTCCCGATCCCACTGCCAATCCCCGAGGGAGGAGGAGG
CAGCCACATACATGAGCCAGGATAAATGGCGCTTAAATATTACAGTTCGAGTGTGACAGCAATGGATCGGCCAGTTGGGTCACAATCAGATGTTGGCTAGCTCCGCGACGA
GGCCCGCTAGGCCCATCATATCTCGGCATATCTCCGTGGATTTGACGGGAGCAGGAACAGGCCCACTGTGGTCAATGTTGCCAGTTGCAGCTCGGGTTAACTGGAATTAAG
GAAAGTTCGACATTTAGCAAAAGCTCTTTCATGTTAATAAAAAACGAAGTCAGAGGAAACAAATACCATTTGCATATCCCTTTATGCTGAATCTATTGTGTTTGTATTA
ATTTTATGTATGACGCAATTCATGCTATTCTCTGTGACAAATTTCCGAAAATGAAATGATTTGATTTACAGACATTCCTTCAATCTCATTTAAATTCGAATCTGTGCAAAAT
TCAATTTTCAATTCATCTCGCTGCAATTAAGCAATTCACAACTCAAGGCAAAAAGTGCACAGCATGAATATACCGCTCTGTGTAAATTCGTTGAAAGTTGTCAACGGTGT
AAAAGATTCTTAAGATGGCCTTTGATTTGGGTTGCCATTACCGCGATACAGTTAAATACCTTTACTTCTTAAGTAGATAAACGTTAACTCCTTTAGTTAGTCAATGCTTTCAACG
AAAATGCCGAAGTGCCTGACTGTGAGCAATAAATCAAAATTAAGTTTACCTTATCATCTAGAGACACCCTGGCTTGATTTAAATATGATATGATCCCTTTGCTTTGG
ATTAGGTACCGCAACTAGACGTTAAGAAATAAATATCTGTGGAGGATTTTCTATCAGAAAATTAATAATTTTATACAAATATTCACGGTTGATTTGGTGAAGAAAT
CAAGAAAATCATCAACTTCGGAATTCCTGGCGCATTCATATATATGATAATATGAGCTTTATACAAAGTAAGCACTATTACTACAAAGTGGCGAA
ATGATATTACCAATAAATTTGCAAGGGAATGGCTACATTTTGGGAATATTTTATTT
(SEQ ID NO: 82)

Exon: 6448..5911
Exon: 4171..4069
Exon: 3988..3863
Exon: 3132..2949
Exon: 1945..1001
Start ATG: 6448 (Reverse strand: CAT)

Transcript No. : CT22855
ATGGCTGTCAACTTGCAGCTGAAATAATTAAAGCGCCATTATCCGTCGCTCATGTATGTGGCTGCCCTCCTCCTCCTCCCTGCGGGGATTGGGACTGGGATCGGGAGTGGGAG
TGGGTGGTCTCCGGGAAACAACTATCCATCGTATATGAGGATCGCGGCAATGGGACAGGTGGAGGAGAGT AGGAGAGAGT AGAGAGTGGATATCCACGCGGACCGAACCGGTAC
GGATGCAGTGGCTCTACCGAAACCGGACCCACCGCGCCGTGACCATCTTAATTACTACACAGCATCGCGCGGCCCGCGCAATGGGCGCACCTTCTACGATCTGGTGCTCT
TCTTCGGAGGCGATCATCTGATTCCAGTCTTTGCGACCTTCGCTGTGTCACCGTCTATCGGGAAACACCTGGTCTACTTGGCCATTAGGACATCTCGCGGCTTCGACCA
TTACCAACTGCTGTTGTGATGAGCTGGCGGTGGCGGACCTTCTCGTGGGCATCTTCGTCATGCCCGCCGCTGCGCGTCCATCTCATAGGCTCGTGGCAACTGGGCTGGGT
GCTCTGGCAGATTTGGATCTCCTCTGGACGTGCTCTCTGACGCGGATCATCTTACGCTTGTGGCCGATCAGTGTGGACAGATTGTGGCCGTGACGAGCCGCTACGTTAC
TCCGTAAACCGCGCTCAAACGATTTGGCCCTCATCATGATCCTAATCGCTTGGCTGCTCGGCTTGGCCATCACTCTGCCCCCATGTCTGGGATGGTACGAGCGAGGACGAA
GGGATCTGCGCGAGTGC CGATACAAC CAGAACGAGGGCTACGTCATCTTCTCGGCCATGGGCTCCTTCTTATACCCATGGCAGTCATGATTATGTGTATGCAAGAAATTC
CTGTGTCATTGCTACCA GCGACGACAAATATGACGACATAGTGTTCACCAAGAAGAAATTC AAGCGCTACACGCGACGGCGTGAGAAATGAGCTGCGGACGAGGACG
CACAGCTCGGTGGGCGACGCGCAGAGACAGGCCATCGCAAGCTTCTTCAACACGACGATAGCAAGGAGCTCCAAAGACATGATGTCTGAGGACAGCGCAAAATTTGCGGG
CGATGGGCGCTGGAGGAGCAGGTGTTGGGGTGTGGAGGAGCAAGTAGTGCCATCTGGCGGCGCACACATCTGCAATCTGCTGGCCCTTCCCTTCCGCGGAGTAGGTGGTCCAT
GGGCTCGCGCCAGAAGATGGGTGCTAGCAACTCACAGACCTTCTCGTGAAGCGGCTTCCACGCGCTCAACGACCATTTACCACATGAGCATGGCATGGGCTGGGACGAGT
AGTCTCTGGATGCGCAATGGCAGTCTCAGCCGCGCGGCGCAACCGCCAGTGTGACAGCCATCCTTATCGAGCCCGCGGAACACAGAGCTTAGGCATTTCGATGGGG
AGCGGGACCGCGAGAGATCTGCGAAGCTCATCCACCACTCTTATCATACCCAGCGGGCGCTTACACGACATCGCATCTGCGGGAACACCAAGTGCCTCAACCAAGCATGA
ATCGCTGTCCAAATCGGATTACGCTGCTGAAGAAGGAGAACAGAACCTCAACACCTGAGCATTTGTGGTGGTGGCTTTATCGGCTGCTGGCTGCTTCTTATCAACTAT
CTGATACCCCGTCTCTGGCGAACACCAGGCCAGCCAAATGCTGGCCAGGCCCCTACCTGGCTGGGTTGGTTCAATAGCGCCATTATCCCTTCTATCTACGCCCTTCTACA
CGCTGACTCTCGGCGGCGCTTCTGGCGCCTCACCTGCAACGCTTCTTACGCGCGGCGCAGAAGCCAGCATGCTCCCCACGAAACCATATGTCATCAGGCGATAG
(SEQ ID NO: 83)

Start ATG: 1 (Reverse strand: CAT)

MAVNIQLLNLSAIYPSLMYVAASSSSLRGLGSGVGVGGLPENTSIIVEDAGNQTGGGGVGVGGGGYSPGNPTDAVALTEPGPTAPVTTIFNNYNSAAAAEWAHFYDLVL
SWQGGIILIAVFATFIVTVIGNTLVILAITLTRLRLTITNCFVMSLAVADLLVIGFVMPFAVAVHLIGSWGLGWLCDIWLSDLVLTCTASILCSAISVDRYLVALTRPLTY
KRRRKSKRALILIMILVLLAALITCPMLGVEYEGRRDLRECRINQEGEYI FSAMSGYTFIPMAVMYIVYARI SCVIA SRHNDNDITSVHNKKFRYTAADVENELSEQEQ
HSSDVGQRQQRATSRTFSSQGT IAKELQDMMLSDSNDNCAAMGAGAGGGGGGASATGGTGHQCSLLAL PSGGVGSGMGCAKNGCYELTRPSSLKRASTASTTITMTSGMGPGS
SLLDAAQWGSQPPGQTGQVQTHSLSPQPTHSTFRHSHGEDRERLSHHHHPHYHHQAGVTTTSTSGNTSANTNSKLSNRNITSLKKENKTTQLTSLVVGGFIAICWL PFFNT
LITFLAEAHQSQMLAKALTWLGWFNSAINPFIYAFYSVDFAAFWRILTCKREFSAGQKQPQPTNTMSIRR*
(SEQ ID NO: 84)

Name: Octopamine receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384651

GTGCTCGACATTTCGCTTGGCTCCCGTAAGAAATTTGCCACCACGATGACCTCGTGCCCTGGACCATCAGATCGTGCACCAAGTGGGAGCCACGAAGCCAGCTCCTCCG
GTAATTAGTATGCGTTTGGCATCTTATAGTTTCAGGTATTTAACTTCGGATACCTTGGCCGTGTGCTGGTCTGGAAGCTACGAATTTGTCTCCAATCTGGCCAAATTTCT
CCTTTCGCGCTGGAGTTTCGGCTCTTGTTCTCGAATTAAGAGAGCTCTGCTCCTCGTAGCCCATCTGAAGGCTCCTTTGTAGCGGACTTTCCGCTGGCGGCCCATTTGGCCTG
AACCTCCTTCGACGCGCGGAACGGAAACGGCGACCTTTCCGCTTGGGCAGAACTGGCCATCTGGTAAAGGTAGACACGAGAGCAGGAGGAGGACGACGCGATGGCTGCAACGATC
TTCAGACGCTTTTGGTGGCAGTCATCTCGGTTATGGGTGTGTGCTAGCTACGGTTCCAGTGCTTATCAGCAACTGATATCGCCACTTCGTTCCGCTTCGGCTTTAGAAAT
CATATAAACAATAACAAATTCGGTTATCAGGAGGTGGCAGGATGGCTCAACCGGAATTTCTAAATGGATTATAAATCTAACGCCGAGGGAAGAGCTCCGCCCTAAT
TTTAAATTTTCGATCAAAAAGTATCGACAGGCGAACATTTTGATTTTGGATTATAATTAATGAATGATATGAATTAGACATTTTGTGGTGATCTTCCCGAGGGACG
AATATTTTTCGTTTCAAAAGTAAAAAGGTCCGAAATTTGCCAAAAATGAGTTTGAACACTAAATTTGTTTGGGCGTTTCTAGGCTGGTCAGCAGGGGACTAATTTCCGAATA
AAGTGTTTGCCCAAACGCGATTGAAGGCTATTTAAAGCTGCCAACGATGAGCATGATCGCCAGCTCGGGCAAGTTTCTAGTCCGAAGTTGTGATGCTCAGCATAGAGCT
TCCTTGGGTGATTTTCTGTACAGTTCTGCTTTTAAATTTTACGAAGCACTCAAATGCGATATTCGGGGTGCAACTACGACGACGGTTGATATCTCATACATTTGAAGG
CAAAACGATTCGTATTATATGATGACATCGAAATTCCTGTGACCACTTAATCGGATCTACGAGTTCAGGCGAATTTGGGCGAGTTGGGAGGTTCCGATTTGAAAGCATTTAA
GGGCTGTGTCTGACGCTGCGGCCCTGTATTCGATCTCTGCTGCCAGCAAGAACTTTTGGCCAAACGAAATGCGATGATGGTCTCAAAGAGGAGCTGCCCGGTCA

41/89

Exon: 1001..1673
Exon: 1849..2025
Exon: 2087..2338
Exon: 2403..2836
Start ATG: 1001

ATGAGGCTTCCTTGGGTGATTTCGTGACAGTTCTGCTTTTAATTTTACGAACAACCTCAATGCAGATATTCGCCGGTGCAACTACTACGACACGGTTGATATCTCATACA
TTGAAGGCGAAAAAGTCGTGATTTATATGATGACATCGAATTCCTGCTAGCTTACGAGTACGAGTACGAGGCTTTGGGGACGGTTGATTACGCCGATGAA
GCATTTAAGGGCTGTGTTTGCAGCGTGGCGCCCTGTATTCGAATCTGCTGCCAGCGCAAGAACTTTTGGCCAAACGGAAATCGGATGATGGTCTCAAGAGGAGGCTCGCC
CGGTTCAACCCCTATATATACTTCCATACATACATGGACCTACAGGCAGAGTACCCTACCAGATATGGCTATTATCAGAGATGAGTCTCTTGATTGTGATGAAATGATTTACA
TCAGCGACTTTTAACATTTCTTAGAAGAGGTTAGCATTTCAAATCTCAATAAGTGGACCTATAGTTTGGTTCCAGGATGGAAAATTTGGGTTACCGTTGACCTCTTCAT
GGAAAACAGCAGTACTGCTTGTACCGCGCAACCTTTGATTCCGATTTTCCAAAGTCCATGTGGATAATACGACACCGCTGCACAAAGCCACATATCTCTGGATCTCTTAGAG
ATTCTAATTATAACAATGATATGCTTTGTCTCAACAATCGCAGATATATCTATACATTAAGAACTGGCGAAATGTTACTGGCAAGTGCAATTTGATGCTGTATAGTTTCGAGGT
TATTCAGTCTGCTGATCATGATACATGATCATTTAAATCTAATGGAATGGCATTTTGTCTCCAGCTGGTTACAGCTCGCATCTTTTCCGATGGCTTCCAACTCTGGCTCTC
CGTACACGCTACCATACCTGGAAGTCTTGACGTCGCTCAATCGAGTCAACCTTACTCGGTTCTCGGTTACACAGCCCTTCTGCTGGAGACACGCCGCAATCTAGACG
GGAAGTATTTATAGTTAATCACTGTTTGGGAAACGATCCAGTAAATGGAATGGTTGCCCTGGCTGGTTTATTTCGGTGTCTCGGTCAAAGATGGCACCCATCCGCTCT
GGATCTATATAAGTGGACCTCGCTGGCCCTCGAGCATTTCAATGTCGCCATGTTGCCCTGCACCTGCACGCTTTTACATAGGAAAGTGAAGGGGGTATTAATAGTTACACA
TAGGAGGAGGAGGAGGATTAATCGCATAAACTTTGACAGCCAGCATAGATATGATGAAACCTTGGACCTTGGTAAACTAGTTATCTTGGCGTATATCCCAATAGTTAC
CTACAGTTCCTGCGGCTCTCCATCGTATGGGCTTACTTGGATATTCAAATGTCATTCCGTATTCCTGCACGGCTCCACATTTTCTGGGAATGGGTCTGGGATAATACCGAGT
ATTTTCCACAGCGCGTTTGGAAATGTTTGTGTCGTTCTGCTCGCTGAAGCGCAGCAGCATGGACTCTAATGATGGATTCT

(SEQ ID NO: 86)

MRLPWVIFCTVLLLIIFTNNSNADIPGCNYYDTVDISYIERQNSDLYDDIEIPASLTGYGYEFRQFGDGSITPIEKHLRACVCSVRPCIRICCPAKNFLANGKCDGDLKEELA
RFPKPYIYFTYMDLQARVPLTMAIIRDEFFDCDEMIYISDFNYFLEEVSIIQIFNKCGLIWVFGDQGFVVTVDLEMEKQDQCYLRHNFSDSDFPKSMWIIHRCTSHISPGSLE
ILIITMTCFVLTIIVLYYIKKLNRVNTGKCIVCCIVSRFIQCLIMILDHLNLLNGICSPAGYSSHFFRMAASNWLWSVISYHTWKVLTSLNRVDPNRYFLRYNAFVWSTAAIMT
GSIYIVNQIWENDPSKWNWLPVGFIRCSVKDWHPSVWIIYISGPSLALSTFNVMFALTAIYIRKVKGGINKFTNEEGRINCINFDSQT*
(SEQ ID NO: 87)

TGAACCTGCACCCACCAGCCAAAGACCCAGCCAAAGACTCCGACCAAGGCAGTGTCTCCAGCCGCTGTTGCCAGTCCCGCAGCTGCGGGCAACGTCCGCAGACGGCTCTCCATCT
 CCGCGCAAGAAGGCAACCGCTGCTCGCGCTCGCCGCGAGGCCAGGCCACGCCGAGCAACCGCGCCGGCTCGCTACAACTGCCTATCAAGTAGAGCTCGGAAGGAATCT
 GTAATTTAACTAACATCAAACTTTAATAACAGAAAAAATTTACATTACGCCCGAGGAGGATGAAGAAGAACAGATCAATTCGATCTTCCGCAATTTGACATTTG
 GCATCCCAATTTGCATATACACTGAATGAATGGAACATGAGAAGAAGACGGTCAACTTTTAGTTGAACAGTTGAATATATTGTACGACGAAGCTTTCTGAAACATGCAGGAGG
 GAAGACCAAAACAAATTAATGAATCGAGAACTGTTATCTACCCGATCCCCCTCAGACAGCATACCATGCCAGCCGCAACCAACCTTCAACTTATGTATCATAT
 ATAAATCTATAATTAAACAAAGAACAAACACTAGTTGTAAACAAACAGCATTCCTGTGTCTATTGCGGATATGAAATTTGGCTTTCAATCTGTGCAAAATGAGATTC
 GTAGCAGTGTTATATCTATGAGGGCTGTACGATTTCTGTTTACGCGCAATTAATACATTTAATACATGTGATCTTTTACACAGCGTTATTTAATGTCAAAATCCGTA
 CAGAGATTAAGTGTAGGTGGAATACATACATATAAGGAAGACCTTTGATATAGTAAGATACGAAGTGGGATATTTATTATTACAGCTTACGAGACTAGGCGCTACTCCCA
 GGAATGCTCTGAGGGAGCGGAACCGCTTTTGTAGTAGATCCCATGATCGGAATCGGGATTAAAGTATCTGGGGGCGCGGGTATGAAGATCGAGTGGAACTAGTTGA
 ATTCCAGAGTATTCGCTCCGTCATCTGTCGTACGCATGCGGCTCTGGTCTGCTCTGGGAACGACGAAGTCCGCCGCGGACGCCATCCGCTCTCCAGCGCTTATTCGATCCCG
 CAAAGGACAGGATTTAGTGACTTGGAGCGCATCAGCATATGACATACGGAATCCGAGGTGAAGTCGACCGGATGAAGCAACATCGGCAGTGATGAAGAAGATTTGTTTCAGTCGGGG
 ACCCGATTTTGGAGCTATAGGCCAACGGGATGGCGATCTAAACCGGAGATAAATGTTGATAAGCCGATGAAATTTATTATATATTAGTATTGCATAATACGGAATGGTT
 GAAGATACTAGACGTTTAGTAGTTTACTATCTCGAAATTTAGAATGGTATCCACCCGATACCTACCATCTGTTGGCATCCAGACATGCGAACGAGATGCTGAGGAACCGCAT
 TAGTTTGGCGAACTTGATCTGACTGGCAGGAGTCCGTGGCGCCGCTATTTGACCGCTAACGTCGTGCTGTATGCTTTGCCGCTGACGCTGATCGATGGCTTCTCCGCT
 CCGGTGCTACGTCGGCTCTGTAGAGGGTGGTGGCGCTGCTGCTCTCGGATCATGCTGATGGCACTGTTTGTGCTCCGGAACAGGCTGTAGTGCATATGCCGCTTG

[illegible]

Exon: 4340..4237
Exon: 4024..3582
Exon: 3513..1001
Start ATG: 4340 (Reverse strand: CAT)

[illegible]

44/89

CACCCATCGAGGTCCTAGCGATCAAGTGCATCAGGGTCTGGTGCAGACCATTTCGCCACCTCCCGACGCTTTCAGTGCCCCACCCCGCTCTCCGATGACTACTCGGATCG
TCCCTTTGGGAATAGTTCCCGTAACAGTGAAGTGGCCCTGACCTATGACCTGATGACGAACCTCGGAGCTACGCTATATGGACGAGAGCTCCGCGATGCTGGCCAGTGTGACG
GCCAATCGACGACCTCCCGAATGACAGCGTCCAGGGGAAACACAGTCTGCTCCACCACCGCCGACGCGGAAATCCACCAAATCGAAGTGAACAGAGCTGCA
GTCAAATCCCGACGCAAAACAGAGTCCAGGCCAAAGCCTGAGCCTCAACCTGAATCCGAATCACAACAGAGTCAAGTCAAGTATCGAGGAGGCGGTGGCCATTGAAAA
ACGTCTTCTCGTCTCGTACAGGAATCGAGGACTTTGCTAAGGTGAGGCGGGAGAGCTGCATTGAGGCCATCTGCTGTTGGATGTGCCGGAAGCTAGCTGCGGACTGT
CCCCATCGCGCGGCTCCAGTCTATGGAGACCAGCAAGGATGCCATCTATTCCCGCATGCCAATGCCACTCTACCCAAAGTAAAGACTTCAACGCCACAAA
CGGGAACGCCCCAGTCCGCGCGGGGTGTAACACACACCGGTGCCACTGGAAAGGATTGAGGAGCGGAAACCTCGGACAGTAATACCAACAAAGATGCCCTCCACCTCGGGCAC
CACCAGCAGTACGCGCGGTGTGGGTGGAGCGGGGAAACAGACGGTGTGGTGCCACCAAAAAAGACGCGGGGATGACGATGCCAACAGGGAGGAGAGAGTTGGCGGCC
AGCCGAACCTCCAGCAAAAGGCATTTCATTTCGATAGGAAAGCACTTCAAGAGCAAGAGGCTCTACCCCTGATCTTGGAGTGGGTGGACGGCAGAGTCCAAGTCGG
AGAATCGTCTCGCAAGCGTTTCGCACGATCTCGTTCATTCTGGGCTGCTTCGTTGGCTGCTGACACCCCTACCATGTGTTGGCCCTCGTCGAGGGGATCTCGCCGCAATCC
GCCCTGCATCAATGAACACCTCTACATGTTCTGCTACCTCTGCTACGCGCAACAGCCGATGAATCTTTTGTACGCTCTGCCAATCAGCAGTTCAAGAGACCTTT
ACGCGGATCCTCAAGGGGATCTGCACATGACCTAA
(SEQ ID NO: 92)

Start ATG: 1 (Reverse strand: CAT)

MRSLNKSAGEEELTGLNGNSSDIYTIKNKWITDMSLRERDRKLSFIFKRVLTHTECKRLLEAAKIYGESSDFNSWPVNEQLQWYRNFVNNNTNLLAVRNQSSDGGSTMS
GSSSDSEILGPVLPFFALWOTILIAICIAIILTVGGNILLVLLAFIVDRNIRQPSNYFIASLAATDMLIGTVSMFFYTIYVLKGYWDLGPMCLDLWLSVDYTVCLVSQYTV
LLITIDRYCSVKIAAKYRSWTRTRVIMVTIWIIPALLFFISIFGWEHFTGKRDLPLPGQCAVQFLKDPINFNTALIIIGYWTTLIVLFVLYAGIYKTAIDMKRSEAKQRK
MQSMVALSAGAMSGMAGHAGIGVIEEKILKTKVELAGDQTDLSACTTVIKRLSGSQANPLAVATEEVEKMTPEORRASAAKIQEAKKREAAVDAEKSERSSSPAFDSD
EESSVNAQQLITQKLNMRKRSSISGLVFGAQAALLATRGKGNLQKSTTNSKSIEMHQQYHHHQHHHNSFLORAQSKEEMRSLHHHQQQPPHQQHQQNPNPFLDRPK
RTSCSTLSQIAEHDRDLVLSAPPTLLNTSDPLSPVDLAPIEVPDQVHQGLVQTLPLPPDAFQCPTPLSDDYSDRPFNGSSGNSELALTYDLMTNSELRYMDESSAMLSVT
ANSTTSFNDVSQKPPVLPPLPPARRNPFSKSNLQSLQTSQNSQSPQSLSLNLNPNHNSQSQSIEEAVAIEKRLLVSYTGIEDFAKVRRESCEIAICLLDVPGLKLAADC
PHRRASSPMETSSKDAIYSPMPPLSPKVKSTSTPQTGTQSAAGVTPFVPLERIEERETSDSNTNKMFPSTSGTTSSTGGVGGGGEQDQGVATKKNAGDDANREKSLAA
SRNSSKRAFHSIGKHFSSKALPLILGVGRQSKSENRRARKARTISFILGCFVACWTPYHVLALVEGFCRHFPFCINEHLYMFSYFLCYANSPNPFYALANQGFKRTF
TRILKGLHMT*
(SEQ ID NO: 93)

Name: Muscarinic Acetylcholine receptor-like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013384504

GACATACGAGCGGAGAAATGATTGATTCTGTTATTGTTGCTTATCATTATCGATTTCGATTATTCAAAGTCCCTTTTTCGCCCTGCGGCGAACGGCGGATTGCCCTCAGT
TTCGAGTTCCTGCAATCAATGCAATTATGTCGTTCAATCGTCTGAATGCTGAACGGTAAATCGAGTAAACAAGAGTGGGCGGAGGTGGTCTGAATAATTTATGACCGCC
CATTAATTATTACTCATAGAGCTGCTCTATGACTTATCGACTTTAAGCCAAATGGAGAGGGGGAGTGAAGCTGGCAGGGCAACCGATCAAAACGAAAAATCAAGTGTGG
GTATCATTTGTATTCTGTAATTAAGAGGCCAATTTGGTGGCAATTCCTGTTTGTGATACAAATTCCTGGCCGCCATTGTACCCGTGGTACTTCTAGGAATTCATTAATGTG
ATCAATTTGCTGTCAGGTAATTTACTTGGTGAACAAATTAAGTTCAGTAGCTATGTTTAGTCATTTCAAAGTTATGCTTTACAAATTTATTTATATAAATCAATGAGTGC
TATTCCTAAATTAAGCATCTGCACTTAAGGTTTCTCTATAAGGAGCCTAATTAGCATGCGGCATTTCAGCAATCAGAGTAGAGTAGAGCGCACAAATAGCAAGTGAAGA
CGCTTTGGTGGCGTAAATATTTGGCCACCAATATCCGTTTCGTTGGCGTAGAATTCGTCCTGGTGGCGCGGTGAGCTCTCGTCTCTTTCCTTTCGGGAGTCTGTAATTCGCG
TCCTCGCTCATTTCCGGTGGCGGTTTACATTTGCAATTCGAATCGGATGAAATCGGAATCGGAATCAGAATCGGCGCGAGGACATTTCCATTCCTTCCCATTTCTCTT
TTTATTTTTCCTGCGATCTGCTGATATGCGAGGGCTAATTAGCATGCGGCATTTCAGCAATCAGAGTAGAGTAGAGCGCACAAATAGCAAGTGAAGA
GGATGTGATGCTCGCTAGGTGGCTACAACGACAGCGGTGGCGACGATTGGAGCAGCTCGGAGCAGCTGGTCTGTGGGAGGAGGATGAGACGACGACGACGACTGCTAAT
GCCACAGTCGGGACATAACTGATGTCGTCAGGTGGAATGCCACCGGAAATGCGACCATCAGCGCGACCTTCGAGGAGCTACCTTCGATGCGAACAACTACTGGCGCC
TATTGGCCCTCGTCTGCTGCTGGAACAGCGCGCGGAACTCTGCTGCTGCGCATTCGCTGGGAGCGCGGTGCGAGAGTGTGACCAATGCTACTTCTTATGCTGCTG
GGCCATCACCGATCTCATGTCGCGCTGCTGGTTCATGCCGCTGGGCTCTTACGCTGGTGAAGGTGAGTCTCTTGTCCCATTCAAAGTCCGGAACAGAGGTATTTTTCG
AGCCACCTTTTGTGAGGGACTCAATAGGACACTTGAGTAAGTCACTTGTGCGCCAGGATTGGCTTGTGCGGCTAATCTTAGTTGGAAGGTTCCCTTTCCATAATTTTCATAG
TTTACTCTAACGAAATGAGTTGTTGGGAACTTGATGAAATTAATATTAATTAATCTTGGGCGAAGGAAAGAGTGAAGTGTGATAGTTGTTAAATTTTAAAGGCT
TTTAAAGGTGACGAAGAAGATGCAAGGAAACGCAAGATGTAGGGAATATACAAATTAATTTAAATCTTAAGATTCTTATATTTGCAATTTAAATGCGGTAAATGCGAA
TCAATGAAATTAATGACAGGGAATTTGGAAGAAATATGGATATTTACGGATTCCTTTTGTGCAACTCAATCCGAATGATAATCTCTCAAGTGGAGAGATATAATTAATCAA
AGCCAGGACAGAAACCTGGCAAGACATTCAGGCCAGGACCTCTGCGATGTCGCCAACACGAGGCGAGGAGTTGCATTTAATGAAGTGTGCTGGTGGGAATGGCGTGGCGG
TGTGCAGATTGCATTATGCTCAGATATTTTCGATCAAGCTGGCAACGTAAGTAAGCCCGGATAGAAAAATGGAATCGCTGGTGTCTATGGCCGCGCAGGCGGAGCGGAG
CTCCATCGTTAGACAAATTAATAGCGCGCGGCTTTGTAGCCAGCCTGGCCAGGACGAAACCGCTGGCGCAGACAGGGAAGCCAGAAACAACTGCCAGCAAGATGGCAACGG
AAGCTTCCAAAGCCTTTGCGAGCGCGGAGCACTGCAGCATAAAGATTGCATTAGGGGCGGCAACAGAGCGCTGGTGAATTTTCGCTTTCGCTTTCGATGAGTGTGCTTCTT
CGATTTTCGAAACGAATGGCTGATTGAGGACTGCATTATCTGAGTGTGCTTCCGCTCTGCTGCCCTCTTGTGTCGAACCCCTGCAGATGTTGCAACACTGATG
ACAGTTTGTGTTGTGCGCTTTGGGAGCAATTACAAACACAAACACAAACACAGCAGCAGCGGACCAATACAAGCAGAAAGGGGACGACACACAGCAACACACACA
CCTGGAGCAATCCTTGGCAAGTGGCAATGTAGCAAGAGGAGGACCAACAAATGGCTGCTTGTATATCAAGTGTGAAAAATATTGCAATGCAAGTGAACGAGGAGCAATG
TTGCTGTAGTTGCAAGGACACAGCCCGCAACTGGAGGCAAGAAAGGACGAAATCGGGATTTCCTGCTGTCAAGGCTTTGCAAGAGGCTCTGCGACAGAGCTGAATC
TCCAGCGAACAGCAGGCAATAGATCCCAAGGATTAAAGCTCTTACAATTCAAAGGCAACAATGGCGCAAGTAGCAACTGGATATGAGTAAAGCCGTGCAAGTAAATTTGCATC
GGTTACGTAAGTAACATGTGGCATTAAGTGGCATGAAATGGCTATTAATTTCTCTGTTGTAAGTATGCAAAATCAACTTCAATGAAAGCTGAACATAATGATG
TCATGACCAATAGGCGTATGATTAATATTAGTAGACTAAATGTTTAAAAACGCTAAAGGTGTAATAACAAATTAATCTAACAAATCATTCATGGCTTTGGTTGTTGACTT
TCTGGAGGTAAATAATACAAATTTAAATTTTCGTAAGAAACAAATTTGATCGGTAAGTACAGCAACAAATCTAACAAACTTGAATGGAATCAATAGCTGAAATAATA
TTTATAAGTGAATATATATTTTAGGCTTAAGTTACTGAAGCTTACGCGGAATTAAGTTTAAAGTGAATATTAATGCTATTAATGCTATTAATGCTGTTGAGTTTCCCA
GATGCAATATCTCTGATTGTAACCTTCAACGAATTTTCCACTGATCAATCGACCTTTTGGAAACGAAACAGACCCATGCAATTAACGACATGCGTTGATGCCATTT
TAAGCGCTTGCACTTGAGTGAAATGATTACGCGTGGAAACATAATTAATGCAATTAATTTTCAATTTAAATACAGCATTAACAGCCAGTTAAATTAATACAGTAAAC
GCAATTAAGTGAATAGCAATTAAGGCAATTAAGGCAATTAAGTGAATTAAGTGAATTAAGTGAATTAAGTGAATTAAGTGAATTAAGTGAATTAAGTGAATTAAGTGAAT
TTTGAACCTTAATTAATTAATTAATTTGAGGACGAATTAATGATAAACAGAGAACAGCGCAATTAAGGAAAGAAAGGCAAGCGAAAGGCGGAGGACGAGAAAGCTGGAAA
ATTAATGAGCGCAATTTGGTGAAGCTG
(SEQ ID NO: 94)

Exon: 1001..1445
Exon: 1885..2064
Exon: 2784..3172

FIGURE SHEET 44

Transcript No. : CT24036

ATGGAGAGGAGTGTGTATGCTTCGCTAGGTGCCACAAACGACAGCGGTGGCGACGATTGGAGCAGCTCGGAGCACCTGGTCTGTGGGAGGAGGTAGAGACGCAGCGAACGA
 CTGCTAATGCCACCAGTCGGGATATCAACTGCATGTGGCCAGGTGGAATGCCACCGGCAATGCCAGCCTCAGCGAGCCTACCTTCGATTGCGCAACAACCTA
 CTGGGCGCTATTGGCCCTCGCTCTCTGCTGGGAACAGCGCGCGGGAACCTCTTGCTCTGCTGGCCATCGCTGGGAGCGCCGGCTGCAGATGTGACCAACTACTTCCTT
 ATGTGCTCTGGCCATGCCAGTCTCATGTGCTGGCCGCTGCTGGTCACTGGCCATCTTACGCTGGTGAAAGGTGAGTCCTTTGTCCCATTCAAAGTCGGGAACAGAGTGG
 AAGAGTATAATTAATCAAAGCCAGGCACAGAAACGTGGCAAAGACATTCAGGCCAGGACCTCTCGCATGTGCGCAACACGATGAGCAGGTGCATTTAATGAAGCTCGCTGGT
 GGGAAATGGCGCTGGCGGTGTGCAGATTGCATTATGTCTCAGATATTTCGATCAAAGCTGGCAAACTGTGCAACACTGATGACAGCTTTTGTGTGCTCTTTGGGAGCCAT
 TACAACAACAACAACAACAACAACACGACAGCGGCACCAATACAAGCAGAAAGGGCCAGCACACACAGCCAAACACACACTGGAGCAATCTTGGCAAAGTGGGCAT
 GTAGCAAGAAGGAGGACCAACAATGGCTGCTTGTATATCAAGTGTGAAAATATTGCAGTGGCAGTGACAGGAGCAGTTGTGTGTAGTTGCACGGGCACACGCCCCC
 AACTGGAGGACAGAAGAGCAGAAATCGGGAATTGGCTGCTGTCAAGGGCTTTGCGAGAGGTGAAATCTCCAGCGAACAGCAGGCATTAGATCCCAAG
 GATTAA

(SEQ ID NO: 95)

Start ATG: 1

MEEDVYASLGAYNDSSGGDDWSSSEHLVWEEDETQRTTANATSRHNQLHVARNNATGNATISATFEDVPFDANNYWALLALVLVLGTAGNILVCLAIAWERRLQNVNTYFL
MSLAITDLMVAVLVMPGLILTLVKGESFVPFKVRNRVEEYNNQSQQRKRGKDIQARTSCDVANTRHELHMLKLLVGMALAVCLRHHVSDISIKAGKHVATLMTVLVFRFGSH
YKHKHKKHKEHGHQYKQKGRRTHTQTHWSNPWQSGQCSKKEDQQAACYIKCEKYSGSDRSSCCSCSTGTRPPTGGKKGRNRDLPAVKGFAGEPATEAEISSEQQALDPK
D*

(SEO ID NO: 96)

Name: 5-HT2 receptor-like 1

Classification: G protein linked receptor

Celera Sequence No. : 142000013384504

[illegible]

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Exon: 1001..1264
Exon: 1550..1778
Exon: 7511..8268
Start ATG: 1001

ATGAGAAATCGGTTTTTTTTTCTTTTACAGGCTACTTTCTTTGGGCTCGGAGCACTGCCTCACCTGGATCTGCCTGGATGTA

CTCTTCTGCACGGCCAGCATCATGCACC

TGTGCACCATCTCGTGGACAGATATTATCGTACGATACCCAAATCGGATTTGGCCGGATAAAACACGTCGGCGGGTCACTCTCA

AAAAATGTTTTCGTTTGGCTTCTGAG

CATCGCCATGAGCTTGCCTTGAGCTTGATGTACTCTCAAGAAATCAGCCTCGGTCTGGTGAATGGAAC

TTGCCAGATACCGGATCCGGTGTACAAGCTGGTTCGGCTCCAT

GTATGCTTCTACATTTCACTGGGCGTGATGCTGCTGACATATTGCCTGACCGTCCGACTTTTGGCCCCGGCAGCGCCAGAATCTGGGTGGTGGACAGCAGACGGCGGGGCCA

CTCCCGGATGGGCGAGTGGAATGGCTGGCCAGGCACGGCCCTTGGGTGGAGCGCAGGACGGCTTCAGCCACAGTTGGGCTCCACCTTCAGTCCACATCTGCTGCACAAATCA

GGGCAGCTCGCTGACATGCTACAGCGCATGGTAGTGGGTACTTGGCTGCTCGGGAACTCTTGCTGCTGGCCGACAGGAGGTGAGCATCTCAAAGACCGCCTCGGTGGTTCAG

TGGGATCTCAAGTCGTATCTCGCGGGGACAGAGTTTGGCGGAGTGCGAACATCCCTGCTGTTGACGCGCCACCAAAACAGCGACCACCTCCA

ACTCATCCACACCGTTGA

GAACTCGGCCACCTCTGAGGAGTCAACCAAGATTA

GAACTACCCAGCGCGGGCGAATGTGGTGGCAAGACAGGACGACAACATCTCCACTGCATGTTCGACAGTGCAGCA

GACGGTGCAGTCTATCATCTCGCGAACTCCAGCGTGATCTCCAGAACTCTCTCGCGCCAGCGAGGATCTCAGGCTGGAGCAGAA

GGCCACCAAGGTCACAGGTTCTGGAGTGGT

TTCTTTACGTTCTGTAATCCTCTGGTGC

CCATCTCTGCTCCTCAATCTCTCGCCAGCGTGTGCGCGGAGTGCAGGAACGAATTAGCCACTGGGTCTTCGACGTGGTCACT

GGCTGGGCTACGCTAGCTCATGGTGAACCCCATCTTCTACACCATCTTCAACAAGGTTCTCGGCAGCGCTTCAAGAAGGTCCTGCTGTGCCGTATTTCGAGCACGAGTGC

CTGGCGACCGAGCAGATAA

(SEQ ID NO: 98)

MKNRFFSFSTGYFPLGSEHCLTWICLDVLFCTASIMHLCISVDRYLSLRYPMFRGNKTRRRVTLKIVFVWLLSIAMSPLPSLMYSKNHASVLVNGTCQIPDPVYKLVGSI
VCFYIPLGVMLLYCYCLTVRLLARQRQLGGGQQTAAATPGWASGLGAPALGGPAGAAPATVGSTLSPHSAHNGGSSLTVQSDSGYLAAPGTPCPGRRKLSISKTAASVVT
WDSFRHRRRGSFSGGVRTSLLLTPTKTATTSNSSTPLRRSATLRSHQNMNYQGAGCEGGKTRTTTSSPCMLQRQQTVRSHHSRNSSVISRNSSRHGRIIRLEQKATKVLGVV
FFT FVILWSPFFVNLNLTPTVCAECEERISHWVFDVVTLWGYASSMVNPIFYTI FNKVFRAQFKVLLCRYSS TSAWRPSR*
(SEQ ID NO: 99)

[illegible]

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Exon: 7877..7337
Exon: 3829..3532
Exon: 3307..3109
Exon: 2992..2899
Exon: 2753..2579
Exon: 2520..2447
Exon: 2379..2269
Exon: 2040..1852
Exon: 1784..1669
Exon: 1420..1364
Exon: 1300..1093
Exon: 1027..1001
Start ATG: 3796 (Reverse strand: CAT)

Transcript No. : CT24513

CGGTTTCGTTTCGGAATTTCAAAACCTTCGGAACCTATTGTCTCTCAGAGCCGTTAAAGTGAAACTAATTTTAAATAATGTGCGAAGAGAAAATGAAAATTTTAAT
AGCAAAACGCAAAAGAATAACAATAAGTGC CGCGCATTAATGTCGAATAACTTTTC TAGTCGTTGGCTTTGAGTGCTTCGAATAAGTTCAAGGGTCGCTATAACAAATAAT
CCCGCATATAAATGTGAAC TCAATAAAGTTTGGCCACGTAATGGGCGTGTTGTCATAACCAAGGCAACTAGCAAAAATAACCGCTCCAGCAAAAACCGCTTTAACATAAT
CATTTTCGACGCTTGGAGGGCTTAAACGCATTTAGGACGGGTCAAGTAGGTGGTTCGAAAGGACAAAGGACCCGCTCTCTGGCGCTTTTGGCGTCCGAGTCCGAGTGCTAAA
CGCAAGTAATCCCAAGAAATCCACGCAAAATGTGAGTGTGAGTGCGGCAAGGAATCTCATTCTGCAGATAGAAACTTGGCTAGGAGTAGAAGTACAGAGATGCCAAGTGC
CAGGATGTTGTAAGATGAGTGACCACAACCATCGATTTCGTTGACCGCCAGCGGAGTGATCCGCTCTGGATCTGCAACAACCTAGATGGCATCCGTTGAGCGCTGGAGTT
GCAGTGCCTGGTGCAGGAGCACATCGAGGCATCCACCTATGGCAACGATAGTGGCCACTGCTCACCAGTTTCGACTCGATCCTGTGCTGGCCGAGGACAGCTCGCGGTACC
CTGGCGGTGCTCAGTGCAATGGACGAGTTGACGGAATTCATTACGATAGCAGCAAGAAATGCAACGAGGTTCTGCCACGCCAATGGAACCTTGGGAGAAATACCAACTACG
ATGCTTGCGCCACCTGCCCGCCCAAGATCCGTTGCCGAGTTCGAGGTCATCGTGGAGCTGCCACCACTTATCTACTACATTTGGATACACCTCAGCTGGTATCGCTCTC
GCTGGCGCTGATTGTTTTCGCTTACTTCAAGGAGCTGCGTTGCC TGGCGAACCACTTACCCCAACTTGTCTTCACGTACATCATGTGCGCTTTGTTCTGGATACCTCTTG
TTATCCGTCAGATCTCCATCCGAAGTGGAGTGGGACGCTGCAATGCTCTGATCACCCTGTTCCACTTCTTACCCTGACCAACTCTTTTGGATGCTGGTGCAGGGTTGT
ACCTGTACATGCTGGTGGT TAAGACTTCTCCGGGACAAATTTACGCTTTAATATCTACGCTCCATAGGCTGGGGTGGCCCGGCTTGTGTTGTCACCTGGGCTGTGGC
CAAGAGTCTGACGGTCACTTACGACACCCCGAAAAGTACGAAATCAACTGCCCTGGATGCAGGAGACCCATGTGGATTTGGATATACAGGGGACCTGTCTGTGCGGTGCTA
ATAATCAATCTCACATTTCTGCTGCTATCATGTGGGTTCTAATCACAAGCTGCGCTCGGC CAATACAGTGGAGACTCGTCAGTATAGGAAAGCTGCTAAGGCACCTGCTGG
TGCTAATCCCACTTTTGGCATCACCTACCTGGTGGTGGCTGGCTCCCTCGGAATCTGGCCTAATGGGACACATGTTGCTGCTCTCGGAGCGGTATTACTCAGCACCTCA
GGGATTCTCGGTGCTGCTGTTCTACTGTTTCTCACTCGGAGGTGCGAAATGCACTGAGACACCACATTTCACGTTGGCGGACACGCGCACCATTTACGTTAACCAGAAC
CGACGTTATACCAAGAAAGCTTCTCGAAAGGCGGTGGTTCGCCCGGAGCGAGAGTATGCGACCTTTGACACAGCTACTATGGGCGTGGCAAGCGGGAATCGTGCATAAGTT
CGGCCACCAACACGCTGGTGGTGCAGATGCCCACTTTCACTCCACCGGGGCTCACAACATGCCCTGCACACGATGCCCACTTTGGCGGCAATGCCATGATTTCCGG
CAGCACTCTGAGCGTAATGCCCGGCCATTAGTCCCTCTGATGAGGCAAGGACTCGAGGAGAATCGGTGTAG
(SEQ ID NO: 101)

Start ATG: 575 (Reverse strand: CAT)

MSDHNHIDSVNASGSDPLLDLHNLGDIGESVELQCLVQEHIEASTYGNDSGHCLTQFDSILCWPRTRAGTLAVLQCMDELOGIHYDSSKNATRFCHANGTWKEYTNYDACAH
LPAPESVPEFEVIVELPTIIYYIGYTLISLVSLSLALIVFAYFKELRLCLNTIHANLFFTYIMSALFWILLVSQISIRSGVSGSCIALITLFHFFTLNFHFWMLVEGLYLYML
VVKTFSGDNLRFNIYASIGWGGPALFVVTWAVAKSLVTYSTPEKYEINC PWWQETHVDWYQGPVCAVLIINLTLFLRIMVLIITKLSANTVETRQYRKAALKLLVLIPL
FGITYLVVLAPSESGLMGHMFAVLRAVLLSTQGFVSLSFYCFINSEVRNHHISWTRDRTIQLNQNRRTYTKSFSKGGGSPRAESMRPLTSYGRGKRESCVSSATT
TLVGQHAPLSLHRGSNNALHTMPTLANAMSSGSLSVMPRAISPLMRQGLENSV*
(SEQ ID NO: 102)

Name: Diuretic hormone receptor-like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013385198

CATCGATTAAACCACTATCCATTTATGGCTGCCGCTACACAAATCGGCACGTGAACGAGGACTGGTATCGACTTTATAGGACCTACGACACCAGGGATTGTTTTCGG
GCAGTTTACGAGCGCTCCAGCGGTACGAGTTGGACTGAATTAATGCCCTTTTTCAGAAAGAATAAGATTAGACAAAATCTGTGATATCGATCTGTGGTAGCGTTGCAG
AGATCCCAATCGCGTGTGTGGCGAAGGACGACAGACATAACACATTCCTCCGTGCTGAAAATCAATTATAATAGTCTGGGAAACAACTAAAAGAAATATAAAATGGCCC
GTTGGTTTATAAAAAATATAACTTTAATCTGATCTCTTATAATGTAAATCATCATGCAATATAGGAGCAACCGGATGGTCTACATGTATGCTGATATACGTATATTT
TTGTGTGGTGTTCATTCA TAGTGTC AGTGGGTATAGGACTTGTGTGATGTGAGTCTACAATAAACGATAAACAGAAAAGGAAAACCCCAATCAGAAACATAAGAAAATAAG
TAAGAAATAAAAGGATATCTAAAGCCAAACCAACCATGCAGAAATAACA TAACATTAAGAACTGATGTAGGGAATGCACGGTTTAAAGTAGATCACTTTAGTTGGCAGGAGT
GTGTTAGTATATTTAGAGAACGGGTGCAATTTAATTTGGAATTTATTTGGATGGGTGGATGTCAAAGAAGTAGAAAGCATATACATGAGCATTTGCGTATACCGCACCGTAGAC
TTTCCGTTCCGCACTCTCAGCGAGTAGTCTTTTGGAGCGCATGCTTGAATGGAATTAATTTGGTTAAATGACAACAACAACAACAACATCGACGACATAAGCGAACAA
CTACAAATATCAGCATAGCCACATCAACTAGGTAGTGCAATAGTGAAAAGCCAAATATGTGCGACGACGAGGTTATATAAATATAATCAACAAACATATGTGCTAGAAATG
CCAAATGAGTGCACTGCCCTCAATGATTGGTTAGCCCACTGCGATTCTTGATGGAACCTGTTCCGATGGATATTCCTGCTCTCCCGCACCGGGTGAATCCATGCCCTCAG
CGCTGCGGCACCTCTGAGTTAAGGAAACAGTAGAACAGGCCACAAGAAGCCCTAAAGAAATGGAATTTTCGAAGATCAATGTTTATGGAAGACTGTCACTTAATCTATAT
ACCTGCGTGTCTATGAGAAAGGCTCTTATGGCTTCGAAGAGATTACGATGATACCTGTTCCGGGCTGTAGCACCACAGATAGGTGATGCCAAAAGAGAGGTATCAGCA
CCAGCAGCGCTTCGAGGCTTGTAACTATGCCGCTTTCAGGGTATGACGAGAAGTAATTTAGTATGAGTACCTGTACGATATGATATAATAGCATGAAGCTTAACATCA
CTTAAGCCAGAAGTTAACCGGGGACCTACCCACATGATGCTATGAGGAATCATAGATTAAACGAGCAAGCCAGTGATGCAGGTACCTTGAATATCCAGTCAATGTGAGAT
TCACGATCCATGCAACATCAATTTCCAGCTTTTGAATAATCAGAAAGGTTGACGAAGTATAGTATAGTGGCAATTTGGTATCATACCCCAATGAAGTGTGCTGCTCG
AGATGCGGAGCAATGCCCTTGCAATGGACCACACCAAAATGCATACGGCTGGAAGCCAGCCAGGCTAGAGTAATAAGCTAATGTTATCTACGTGAGAAATGTTT
GCACACCAAGCTGTACAGATAGAGGCTATAAAAAAGCAATGATTAALACCGAGGAATTCAAATGTATACCGCTTTTGAACCCACCCCTCACCAAAACATCCAGAAATTTGG
TTAGGTAAAAGTACTGAAACATGATTACCAACGTTATGCAAGCAGGCTGACTGACTCTGTGGTTATCTAAGTGAATAATTAATAGTGGATGGGATATATATTTCCAGC
TTATCTTACCATTGCAAGAACAGTGTGAGTATCCAGAGGAGTCGGATGTGATGAGTGAAGGAAACAAATTTGGCATGAATGGTGTTCAGCGCAACGAAGATCTCTGAAG
AGTTTTRAGCATGAAATGTGCAACTCATGTGTGATACAACTCACTTAAAGCTGAGGAATATGATGAGAGCCACCACCAAGGTGGCGAAGCTCAGGAATTAACCGCG
GCATATATGATGGCCGCGAGTTCCAGCTTGGGTGAGAAGTCGGGCACACCGGATGAGAGCCGAGTTCTGGTGACAGCGGCTCATAGTCCGAAATGATGATCCCACTCCGT
TTGGAAGCAAGGCGGGTGGCATTTGCTGCAAAATATATTTATTTAAGATCTCAACCCCAATCTGGAATACCCACCTGTGGTGTGACAGCCCTTGAATCTTGAATCTT
CGAAACAGGATAGTACAGGACAGCTGCCCGGCTTGTGCTGGCCAGCACAGGACCGAATCGAACGAGCTGGGACATTTGAAATCTGTAAGAAATAACAATGAGAATATA
CAATGATTGGCAAGGTAACCTTCGTTGAGCATTTACTGCAATATAATTCGTTTCCCTCTGTAAGCGCAACTGACCTAGCCCAAAACCATTAATCAGAGTACATAAGA
TTTCACTGCCCGGTTCTGATCCGTTAATGATGCTTCAATTCGGGATCCTGGTCAATTTATAGCTCTGACGATGATTTTCTTAATCAATTTAGTTTCCAGACTTTAC
ACACACCATACCGACGCTAGCTGAGCTGAGCTCAAAAAAATAAAGAAAGAAATCACCAGCTACCGCTTTATTTATATATTTGCAATGGCGTCTAAACAAAT

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TTCGTGCATACGGATACACAAAGTTATATGTACATATATGTATATTAACGTTAAGTATACGACGACCCGGCGATTGTTTACCCAGTTAGTTCGTAGCGCCAAATTAATCGCG
ACGCGCTATGGTAGCGGTGTAGAAACAGAGCTTCATGTATATTTTTTTTATTGACCTTATCAGGTGACGCGGAGCGCGGAGCGGAGACTCGTGAAGAATTACCAAAAT
TGGCAATTCCTTTTATGGCCGGTCTTCCCATTTACTGATAAGCCCCGAGAACCTTTTTCGACAGCTAATGCGATCGGACTTAGCCACGCGACAAGTGAGCTGTCAACGAGTAAG
CCAAACACTGGAGACTGGAACTCCGGCAATATTGTTCTTCTGCTCTTCGATTTCACCTATTAGCCTCGTCATTGGACACAGTCATAGTACGAATACATTTTATGACA
TTTTCGAGGAACACGCCCTTGATCGTGCATTCGGTATAAACAATAATGGCTAAATGCAGCCTCAAGTGGATTTCCTGGGTCTAGTTTGTGTACAGCGGAGAGGAATACA
TAAATAAATATGTCTGCAATTTCCGGAAATAAATGCCAAATAATTTAATCCGAAAGCATCGACAGTTTACTTGCTGGTGTTCATCGTATCGTACTTAAAGTGAAG
AGATTGACGGCGGATGGATTGAAGCCAATATTTCGGAACAGAGATGGATCAATTAGCGGGTGCATTAAATGGGAAATTCCTCAATCGGCAAAAGAGATGAATTCATTG
GCCAAACAAGGGTAATTAGGGAATATTTATTGGC
(SEQ ID NO: 103)

Exon: 2731..2708
Exon: 2549..2432
Exon: 2379..2178
Exon: 2122..2026
Exon: 1971..1878
Exon: 1819..1659
Exon: 1597..1487
Exon: 1420..1235
Exon: 1174..1001
Start ATG: 2731 (Reverse strand: CAT)

Transcript No. : CT24959
ATGAAGGCATCATTATTATACCAATTTTACAATGTCCAGCTCGTTCGATTCCGGTCTGTGCTGGCCACGCACAACGCCGGCAGTCTGGCTGTACTACCTGTTTCGAGG
AATTCAAGGGCGTGCATACGACACACAGACATGCCACCCGCTTTTGTCTTCCAAACGGAACTGGGATCATTTCGGACTATGACCGCTGTACCCAGAACTCGGGCTC
CATACCGGTGGTGGCGACTTCTACCCAACTGCACTGCCGGCCATCATATATGCCGGCGTTATTTCCTGAGCTTCGCCACCTTGGTGGTGGCTCTCATCATATTCCTC
AGCTTTAAAGATCTTCGTGCTGCGAAACCACTTCATGCCAATTTGTTCTCACCTACATCATCCGACTCCTCTGGATATCTACACTGTTCCTGCAAGTGATAACCA
CAGAGTCTAGTACAGGCTGGCTGATACGTTGGTAATCATGTTTCAGTACTTTTACCTAACCACTTTTCTGGATGTTTGGAGGGGCTCTATCTGTACACGCTGGTGGT
GCCAACTTCTCCAGTGATAACATTAGCTTTATATCTACGCCCTCATCGGCTGGGGCTGCCAGCGTATGCAATTTGGTGTGGTCCATTGGCAAGGCAATTTGCTCCGCT
CTCGAGAACGAGCACTTCAATGGGCTGGAATGATTGTGCATGGATCGTGAATCTCACATTGACTGGATATTCAGGTACCTGCATCACGGCTTTGCTGGTGAATCTAG
TATTCCTCATACGCATCATGTGGGTACTCATCTAAATACGTTCTGCTCATACCTGGAAACGGCGAGTATTACAAGGCTCGAAGGCGCTGCTGGTGTGATACCTCT
CTTTGGCATCACCTATCTGTTGGTGCTAACAGGCCCGGAACAGGGTATCAGCTGATCTCTTCGAGGCCATAAGAGCCTTCTCATAAGCACGAGGGCTTCTTGTGGCT
CTGTCTACTGTTCTTAACTCAGAGGTGCGCCAGACGCTGAGGCATGGATTACCCGGTGGCGGAGAGCAGGAATATCCATCGGAACAGTTCATCAAGAATCGCAGGT
GGCTAACCAATCATTTAGGGGGCGATGCACTCATTTGGCATTCTAG
(SEQ ID NO: 104)

Start ATG: 1 (Reverse strand: CAT)

MKASLLYRILQCPSSFDVLCWPRNTAGSLAVLPCFEFKGVHYDITDNATRFCEPNGTWDHYSYDRCHQNSGSIPIVVPDFSPNVELPAIYAGGYFLSFATLVVALIIFL
SFKDLRLRNTIHANFLTYITALLWILFLQVITTESSQAGCTILVIMFQYFLYTNFFWMEVGLYLYTLVQVTFSSDNISFIYALIGWGPCPAVCLVWSIAKAFAPHA
LENEHENGLEIDCAWMRESHDIWIKVPASLALLVNLVFLIRIMVWLITKLRSHTLETQRYYKASKALLVLIPLEGITYLLVLTGPEQGISRNLFEAIRAFLISTQGFVFA
LFYCFNLSEVRQTLRHGFTFRWRESRNIHRNSSIKNRWANQSLRGDCTHLAF*
(SEQ ID NO: 105)

Name: Diuretic hormone receptor-like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013385192
TTTCCGCTGCCATCTGCAGCTGTAGCTGCCAATAAATGCAATTAATAATCTTTGCGCAAGAAATTTGTTGACGAGGACGACGAGAGATACTCGTACTGTATGTGTCGAATA
GTTTGGTGGCAGCAACGGGGCGGAGATAACATATTGCCACGCTCCGGCTGAAATTTGACGGCAAGGCGAACTCAAAAAAAAAAAAAAAAAAGGAATCCCTCAAAAGGA
GAACCAAGAAATGGGATTTATCTCTGGCCAGCTCCAGCCATACAAATCTAATTTAGACCACAGCGGCTTCAGATATTTTATTTCCCACTCAGTTGCGGTCCATCC
TGATCTCTCCAGCTTGCTTGTATGAGTGGCCAGCCGTTGGCCAAATTTATTTAGTCACTGAGGAGATTTGGTTTTCAGGGTTTGGTTCCTGCGGCTC
AACAATGCCTGTGATTTTTTTTTTTTATTTTATTTTTCGATTGCATCCATTAGAGAAAGCACTCTTTGGCATTCTACTAGCTACTTGCACATAAATATCTACGGAGG
GTAGCTTAATAGTCTTATCTCAACTTCCCACTTTATCTCTCGCCGAGTGCATTTTCTGCTGCTAATTAAGTTGCAAGGCGAGCTTCTCAGCAGGCGTTCACGCAAGTA
GATAAAGGAAAGCCCCGATTTCGAGACACTATGCTGGGAAGATATATAATAGTTAAGTTTGGCCAGCATTCGCTAATCGGCTTAGGGGAGCCACCATGCGACCAAA
CAACAACAACAGCAGCAGCAGTGTGCTCCATGCAATATTTAATCACATTAGTTGATGGGGTAACGGCCATGTACGCAACAATTTACAATTCGATTAAATATACGTCTA
GTTGGCAATAGGCGCTCTAATCTATCCCACTTAAATGTATCTCTATCTCTATCTTATTGTCAGGGGACACCCGTTGATAATAAATTAATATGCATCTGCTGGGAAGGCG
GCACTGCACGGCTGCTTGGCAGCAAAATCCAGTTGCAACATCGGTGAGTGACGGGCGCAATCACAAAGGACTTGCAACAGTTGCCGTTGCTTCTGCTATTGCTGTGCTG
TTGCTGTTGCAACAATGGCCAGTGGCAACAATGAACTGAGCGGCTCTACTGCGGACGCGGATGGATAATTTTCATACAAGGTAAAGGACTTTCACCAATGTCACGATCAA
GGCCAAGGGGCTTTTCCACAGACATGGCGACAACATACCAAGCCAACAGTTGGGGGCGTGGTGGGGGCGGAGGCGATGGTGGCAACATCATAGCCATTAGCTGCTGG
CGCAAGGGAAACCGTTCAAAAATCGATTATCGCCCAATTCGGGGGAGCTTCTATTGATTGCGCGTACAAATTTCTCGGGCGATTAAACGACGAGGAGCAAGCAAAACAAA
AAACAGATTTGTCACAGCAAGGTCAACAATGATGGCTGAAATCAATTTAATGACCATATCTACGGGCGCTCCAAAGTGGCCATCTGCTGCACCTATAAAAAAGTGAATC
CGGCTGCGGATTATTTATATATCTGTTGATGTCAGGCGGCTGTAACAACTCGAGATGATGATTAAAGCGGCGCTTAAATGCGGTTTAGGAAATTAATTCCTG
TAATTTAAGCCAGTCACTTGTGAGGTTCTTACATGTAAGCGGATATAAATAGTAACTGCTGCTCGGTGCACTGCAATTTGGCCAAATAAATCTTATGTTAGTACT
TTGATTAAAGTGACAATGCTGATTCTGTAGAGGAATCTAGTTCTAGCTCTCCCAAAAGCTATTAGTTACTCTTGAATAAATATGTTACTTTCTTTTGGCAAAACCAA
CAGAAATTTAAATTTAATATTTGGATTTTTCGCAATAAATGTAATGATGAGGCAACAAATATGCTAGTTTATTTAGGAGCTCTTGGTTTCAATAATTAAGAACA
TAATCCAATCGGCATATAAATCATGATAGCAATTTATTTCCGTGATGAACTGCTGCTCGGTGCACTGCAATTTAGTCAATTAACCTTACCTATTACGAGTATAGACT
TCAACATTTAGACAGGACTTTTAGTTTAAATAGACCCAGCCTGGCCAAAGCAGTGTAAATGACCAAAAGTGGCTGGCCACAGGATCAGCATCCCAAGATGCGATGCCG
CATTTGCTTTAATTAAGGTAGTGTGGATTTGAAAGATGACTGTATGGCAATTAGATGTGTAGCCAGAACCTTGGCCATTTACTTTTGTGTCAAAGTCTGTCGCAAAAT
GCCAGGGAGCGACTTGAACGCTGTACGCGCCAGACAGCAGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCA
CACTAGGACACAGAAGCCAGAAGCTCGAAAAAAGTATAAATTCAAACGGGTATACAAATTCATGTGTCATCGAAGAGAAGCCCTTCTCTGGCGGCAAGGTGACG
TATACGCATATTAATGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCA
GTTTCGCTCTCCGTTGCCAAAGAGAAAGCAAGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCAAGGCA
ATTTAATTTACCAATGCATGCTGCTTTTTCGGCGCGTGTACGCCCGAACAATAAACAATAAGGCACACATCGTCTGGCATTTCATGTTAAAGGCTTCTCATGCTGAT
GGACAGGATACAGAAATATATACGAAGTGAGGTGTGTCTACGTGCCACAAAACAGCGAAATGAAACGCTTTCGAATGCCGGCGACACTTTTGGCGGCACAAAC
CTCGTCTTGTGACCACTCCCGGGGAATTTACAACCTCATCAAACTCAAAAACCTTTCGCTGTTGCTGCGGATTTGGGTCTGCTGCGGAGATTACCTGGGTAA

FIGURE SHEET 49

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Name: G-protein coupled receptor-like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013384565

[illegible]

Exon: 4251..3780
Exon: 3011..2800
Exon: 2707..2212
Exon: 2145..1001
Start ATG: 4251 (Reverse strand: CAT)

Transcript No. : CT27563

[illegible]

52/89

AGGCCAATGGGCAACAATCAGGATCTCCCATAGATGAACAGTCCAGGGAGAGACCGCTCTCCACTCAAACATCTCCACAAATGGCCCACTCCGGCTCTTCTGAGGGCCA
AACTGCTTGGCAACTCCAATAGCGCTCACTGCTGCGAGGATCCACGGCGAGTCTGCTCCCGAGGACCAATCCGGTATCTTTGTGATCGATAGTGAGGGCGAGTCCCGG
CTCAATGGGCAACAAGCCTAAGTATCGAAGGGCACGGCATTCACTCGGAGTTCGCTGAAGAAGAGCCGATCTGCAACTGTAGTCCATCGCTAAGGACGAGGGGTCAC
GACGAGCCAGCAAGTAATCTCTGAGGATCAGGAGTCTCTGTACTTCCACAGCATCCGAGCGAGCAACCATCCACAGAGAACTTTTTAGTCCGCTGAGATCGGTGG
GCAGTTTATGACAGATCTCAACTTGTTCACCTTCTCCAGCCTCAGCGCCGCGTCCACCTCATGACGGCGTCTGTCACGGCTCCACGCCCCTCATCACCACCGCC
CATGGGCCAGGCTCAGGAGGAATCGGTCCAGTGGGTCTGACCACCTCTCGCGCTCCCTTCTGGCGACGCGCGGAGAGTTGA
(SEQ ID NO: 110)

Start ATG: 1 (Reverse strand: CAT)

MSAFRYFSITDPSFEGPLLPLHAATTSKDAKDSPPSELNIPYTVFEVLVAIVSIIIGNVLVIIIFRRERKLRRRTNYYIVSLAMADLLVGLGIPFALLASMLPRNLHAC
LFTVSLLVLCITISIFCLVAVSVDRYWAILYPMAYSRNVTRTAIFIISMCWVAGTIVGLFPLFGWHADVNHNQECLEFVMDYNYLVFLYFATITPALLMLAFYTHIYRV
IIKQVRQIVTMNPASDLSSRSSAAVVQVTPGRGGHTGMLRVLGAARKROVKATQNLIIIVLFMCIWIPLYTINCIAKFCPCYVHPKLTLCIILSHLNSAVNPVLYAY
HLKDFRAALKNLLKMMGVDDIQQAEIHRFSVASQHLQSMDSNMRTQRLVVGESYPIWLROQQEALKNSQLLPKGVVSPCFNNINQTVAAVASVTTDLEREMWNIVE
ASSGAELGETSYEFPSPAPGSRSSERNSSSTVPPAPPAPAKPSVPSASYDNHNSFSQDEDEDDLEFEDVFPASSVPNPVQPIDVELRRSLALVMREKLRSDDTDS
RPMGNNDLPIDEQSRERPLSTQTSPTNGPLPALLRAKLLAGNSNSAHLPGSTASAPAQEQSGIFVIDSEASPGSNHKKPKYRKGTAFTRSSLKKSRSCNCSSIAKGRGVH
DEPSSNLCDQESSVLPHQHPQANHTENFFSPLRSVGSFMQHSNLFHFLQPHARPSTSSASSTASTPTSPPPMQAQEESVPVGLTSSPSLLATSAES*
(SEQ ID NO: 111)

Name: Adenosine receptor like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013384520

GAGCCAATCTTTTTGGGCTTACGTGACGTTGGCAGAACAATCGTTTTGTTTTAGCCATCTGACTTGTGTGCTCTTATGGCCCGCTGCGCTTTGGACTTTGACGCAT
TTGCGTCCGCTTAAATGCGATTGCTGGTGTCTATTACGGCCAATCCTTACTAGTACTTGCACTTCTGTGAATTCGCTCTGGGTTTCCATTTGGCTGATTGCAATTTTCATAA
CAATTTGGAGGATTACTAATGCGCCACGTTGCTCCCTGCATTGCGATGACCTTGAACCACTAGCCACCCTATTAGAATTTCAATTTGTAGTCAACAAGGCGCAGACGACGGGC
CACATGCTAATAAACCACCAATGTATTTTATGGCCAAATTACATTTTTACTAGAATGAATCCATAGAAGATGCGGAAAATAGCGGGCGGGAATATGCTAAGCTTTATTT
GCAGCCATGGGAATATCTTAATACAGCAAAATTCATCCTTTAAGGTATCTCTAGAAGTAACTTTAGTAAGCAACTTTTGCTCGCTCATTTCCATTGAGTTGAATGCTACATTT
TCGAATCTTATTTAAATGCGATTCAATACAATATACAGAATCTACTATGTTTAAAGATCGCAACATTTTTTTTACACTTTCTTAACTAAACACTTTAAAGCACATCTGT
GTTACACTTTGGTATGTCATGTGAACATTTGGTTAACTAAACATATATCTACGCAATTTAAAGGCGGATCTATTAAATTTCAAGGGGCAAGCTTTTCGTTTTTAAATTTAA
TTCCTATATTTAAATGGTGGGCTTCACTAGTATTGATTATGCTTACCATTATCGATAATATTATTAAGTTATATCCGCCCTCAGGCTTGTTAAGAGTAATCCCGCGAG
CTCTGACCAAGTTTAAATTTAAGAGCGCCAGTATGCTGAATAGTATGAGTCAAAAGATTGAATATCTTACACTGAATAATCTGAATACTTGAAGACTAGTTTAGGAGT
TTTTCCAGCCGTTGTCGCGAGCTGAATGAATGAAAACCTTAATGCTTAAAGGAGAGTGTGCGCTCGACTGTGATTAACTTTTAAATGGCCAAATGCAATGAATGCAAAAGCA
GTTTAAACAAAACGCTTTGGTAGCAGCAGCAATAGCAATAGACGACAAATAACTGGCTTAAACAGAAATTCAAATAAATAGCATCAGGAAGCGCACGAGAGAGTGAATTCG
AGTTTATCAGTTAGTTGACTTGGACACCGATCATGGCGTACTGAGCGGCGACTGGCGGCTTGTTCGTCGGATTCTGGAATGCGGATACACATAGAGAGAGAGAGAG
AGAGAGCCAGTTGAATGCAATGAAACCTTAATGCTTAAAGGAGAGTGTGCGCTCGACTGTGATTAACTTTTAAATGGCCAAATGCAATGAATGCAAAAGCA
GTTTAAACAAAACGCTTTGGTAGCAGCAGCAATAGCAATAGACGACAAATAACTGGCTTAAACAGAAATTCAAATAAATAGCATCAGGAAGCGCACGAGAGAGTGAATTCG
AGTTTATCAGTTAGTTGACTTGGACACCGATCATGGCGTACTGAGCGGCGACTGGCGGCTTGTTCGTCGGATTCTGGAATGCGGATACACATAGAGAGAGAGAGAG
AGAGAGCCAGTTGAATGCAATGAAACCTTAATGCTTAAAGGAGAGTGTGCGCTCGACTGTGATTAACTTTTAAATGGCCAAATGCAATGAATGCAAAAGCA
AATTTGTTAGTGCCTCTCAGCGCGGCGGATTCGATGTTGTTTTCGGAATTCAGCGAACCTTTCGATACCTTCTTGCACAAACAGAACGGCTTAAAGGAGTGGGAATCGGT
AAGGTAATTAGTTAAATGAAAAAATTCAGACATTCGAATTTATTAATAAATATATTAATAATATATAAATATTATAAATCAGATAAATTTGAATATTAGTATTGCGCAT
ACTATAACCCCGCTTACCTTCCGTTTACTACCTTGAATGCTTCCGGAATCTGCGGAATCTGCGGAATCTATATATAACAGCGGGTTGATGAGGAGTGGACAGTAGTAGAGGACCGG
AGACATAGGTATCACCCTGATGACAACTCGTCTGATCCCGCAGTTTGGCCCCCGTGCAGGGGCGTAGATGGCAATCAGTCGCTGGGCGTGGAGGGGGCGCCAGCAGAG
GAAGAAGCACACCACCGCCACTAAAGGGGAGGGGGGGGGGCTAAAGTCACTGACGTTGGGATGAAATGGAATGGCAATGGGAATGGGGAAGCACAGCAGGAGTCAAC
AGATAAAGAGTCAACGATTATATTTTTTGGTAATCGCAATGTCATCATATAATGGAGAGTGCCAAAATGACAAACGAGTAAATGGCTGCGAGCGGGTTGATTAGCCGAGC
TTAGGGCATATGTTCAACAGTTGGCATACACTGACGAAAAGTTGAAGTAAAGTTCTTAAATGGTTCTTCAATGATATATAAAGAAAGATTAGATATCATATTTTCAAGATT
TTTAATTTGCAAGACTTTGTGAGAAGACTAAACCTTATTAATAACCGAGTCTTCTACTTTTGTTTTGTGCTGCTGTTTTCGCAAAATTTCTTACTAATTTAATCATCTGAATCTATTAGTC
ATGTTTACTTAATGTTAGCGTTTACGCTTTAGAGATCTGTCATTTAATTTGTGATACCCCCGACTATTGCTGGTATTTTTCGCGAGTGTATATCGTGCTATAATTACCTAGCATCC
TGAGTACTCGCGGGTGCCATAGTATTGAGCCGACCTCTCACCAGAGCTAAGTTGAGCCCCGAGCGGCCCATCAAGCCCGCTGCTCCCTGCTCCACTTCTCCGCGGTTGAA
ACTCATAGCGGTACCGGATCCACCATAGCGATATAGGATCGTATCACTGGGCACACTCTTCACTGCTGCGCTGCGCAACCGAGGACGCTCCACCAAAAGTGGATGGA
TACAGGTGACACCGATAGTAGTACCAACTAGGATATAGACATCGGAGCCAGGAAGAATATGAACGTTGGACAGCTGGAATGAGTGCTTCACTATGACCCGCACTATGC
CGCATTTGCTCACTCCGGAATAGTGTCTGATTCCAAATGGGCGAGCTGCGGAATGGCGTAATCTGAGCATATCAAAACAGGACGATGCGGAATGGCGGCACTGAG
TTTACTCATGGCTGGCCAGCAAGCGGATGGCAATGCAATATACCGCTACCGCTGACCGGCGTAATCGTTAGCACCCTGGCATCTCCGCGCAACAGAGCA
GCTCCGATGAGATGTAATCCCAACACGCTACCGGATCTTGGACAGATGATAGGACACCTCTGCGGAACGCCGCAACAGGAGCAGGAATCCGAGATGGCCAGCGAAA
AGAGGTAGTAATTCGTGGCGGTGTGCATTGAACGGTTCTTCTTAATCAATGAGTACTATGTGCGCCACTACACCGGTTATGAAATAGGAGTAGACACCGTTAC
GGGTATCAGATGGCCAGCGGATCGCTGGCGGTCCAAGATCAGGCTATGCTTCAATGAGTACTGATCTGAGCTGGCAGCAACGCAACAAATACCGAGGTCGAGATCCGTATA
CCGATTACGATTACGATTCCGATTCCCGGCTGTTTTCTAGGGGGCCGACTTCTTCAATAAGTTATCGAGTTAATTTGATTTTCGATTTCGAATTTCTGATTCCCGGAG
AGCAGCTGCAACCGCACTCCAGCGCGCTCCAACTTATCTGATAAGCGCGGCGACATTTTCAATCAGATTGTATCTGTATCTGAAAGCCAGCGACGAGCGTCTGCTTTT
TTCCTTCAATGAACGCGGAAGGTGAGCGCATGTTTGAATGACCGAGCGTCCAGCGAACTGCGCGCTCAATTAATCTTGAAAGCGGCGATATCTTGATGGGACTAGC
CGATGTTAGAGACTAAGCAATGGCTCCGAATACACGCAATTTTGGGTAATTAATAAGGCTAATTAACAAAACCTGTAATAATGTTTGGTTCGGCCATTTTAAAC
ACTTAGTTGGGAATTTTGAATTTGTGATGCATATAATCTTAAATAAACACGATAATGTAACAAACACATCTTGAATTAAGCCAAATAAGATGCAATAATATTAGATG
ATGCAATAATACCAATGACCGAAGGCCACCGTGCATGATGATGATGTCAGGCGCTTACTTGTAGCTTATAAGTAAGAATATTGTTCTTGTATTCTTACTAAAGTAT
ATTGAATGTAACAAATAATAAATTACAATTCAAATCTAATTTAGATTAATTA
(SEQ ID NO: 112)

Exon: 3314..2456
Exon: 1927..1710
Exon: 1547..1367
Exon: 1029..1001
Start ATG: 3314 (Reverse strand: CAT)

Transcript No. : CT27924

ATGTCGCTGGCAATATGAGCCATGATCTTGGACCGCCACCGGATCCGCTGGCCATCGTGATACCCGTAACGGTGGTCTACTCCCTCATTTTCATAACCGGTGATGAGGGCA
ACATAAGTACCTGCATTGTGATTAGAGAAACCGTTCAATGCACACGCGCCAGCAATTAACCTCTTTTCGCTGGCCATCTCGGATTTCTGCTCTGTTGTGCGGGCTTCC
GCGAGAGGTGCTTACATCTGGTCCAAGTACCGGTACGTTTGGGAGATCATCTGCATCGGAGTGGCTGTTGGCGGAGACATCGGCGAATGCCACGGTGCTACAGTAT
ACGGCTTACCGGTGAGCGGATATGCTTGCATTCGTTCTGGGCGAGCCATGAGTAACTCAGTCGCGCCATTCGCATCATCTGCTGTTGGATTATGGCCA

FIGURE SHEET 52

GCACCTCTATTTTGGGTACTTTATAGAAATACAAATCCCATTAAACACATAAACCGAAAATCACATTTAAATTTGATGTAGGTAAATATCCATTACCAACAATTTTTTCGACAAG
ATGGCGGTAGGCGCATGCCAATTTCTAATGCGCAACATGAGCGCTCGTGGCCGGAAATTAATATGATGCGGGATATGCTGAAATGGTGTCCCTCGGCCTGTGGGTTTC
AAAAATCGGGCTGGCTACCTTTCTATGGTGAATAAGCGAACTAATGCTAATTTGTGTGGCTTGTGTAATTTATGCGTTAAAGTCGGATCTCATTACTAGCCACACAG
TGGCTATGGGCAATTTTGCATGTATGCATTTTCATATCCGCGAAAAAAAATCTCGTAATTACCGGCATTATAAATTAAGCTTAACTGTTGCAACTCACCTGAAATGGTATG
CAGAAAGAGGCCAATAATCACATCGGCAAAAGGCCAAATTAGCGATATACATATTTGTACGGTCCGCATTTGCTGGTGGTGGGCAACCCAGCATGACCAAGTGTGCCAA
TGACGGCCACGATACCTAGTTCCCCATAGAAATGGACACAGGCGCAATCTCAGCGGGAGCGGATATAGACGCTTGTGGTAATGAATGAAAAATAGTTGTCATTAGA
TTTATATCACCTCGAAAAAGTGCTTGAACATTTATATAAACTAAATGAGATATAACAACTCTGTTTTACACAAAAATTTAAGAAATGTATCCAATACATAATGCGCTCCTAC
TAAGTTACTGAAATTTTAGTTATTTATACAGACCATGCTTTATAGTGAATAAATTTTGTGTGGTGCTTAGCTTCTTCCAATCCATTAGTTTCAAGTGTGAACTTGT
GAAAAATCGCGCTTGCTGACAGAAAGCATTAATCTAAGTGTCAAGTTGTCACTCAATGTAGCGGACAGACAGACATCTCACGTTCTTGTAGTTTCAACCTCAAGGA
TTGAACGCTATTTTATGCGCAAGAGGGAATTTTATAAATGCCATTTTGGACAGTATACCTGTATATATATACCTATAATCACTAATTTAATAATAAAATTTGATAGTTAA
ACCCAACTTTGATATGAATTTTATTAGCATATCCGAACATCACTGTTTCTCATATCAAAAGTCAATGAGATGGTTGAGGCCAATGTTGGTCTTTAAATTCGGCACCCCCCA
CACTCATGATGAATTTCAAGATCAAGAACTACAGCGGAACTGTCCAGAAGAAAAAACACTTGTCAATGTTTTTGGATGTATGCTCAAGTCAATTTAATCCAGTTGGATG
AGTCCGCTCCCAAGTCTTGAAGGAATAGAGATATACCTACAAATCTCGCTTCTCGCTCCGGCGAGGAATTCAGCGGGAGTCTGCTCATTAAGTCCATT
GCCATAACACTTTTTCTGTTTTTAGTTTTAGTTGTGTATCCTAATGAGGATATGATATCCTTCAAACCTTGTATTTTAGTTGCAATGCTGCTAGCTTATGTAGTGAATACA
TCTTCAGGGGCTCTGAGCTTCATTGCAATCTAACTGGAAGAATATAAAATTCACAGATTTGTTCTATGTTTTCTGAAAAATTCGCAAAAAAGTCTATGCCACAAGTCCAG
TGGTCAGGGAACAACAAAGTGCAGACAACAGAAACAATGATCTGAAAAAAATCAATTGAAAGCTATACAGCAATATCACTGCTTGCCTCAAGGAATGACCAAT
TCAGTGGCGGGACATGAGTTATGGCTCAGGTGGAATCACCTGTGAGCTTTCACAAGTAAAAAGTTTGTCTGTATAATTTGAAAGTGCACCCAAATTCAGTTTGGCGACTGAG
GGCACGTCATAGAAATTCGCTGTATTTTCGTTCTATGAAACAGTTCATGTTTTCATGTTTATGGAAGAGTCACCGCAACGCTCAAGTGTGCTTACGCTTACGCTTATGTA
GGTGGCTCTCAAAAAATAACAAGAAAGTCAGCAAGTTCATTGAATGTTTGAGATAGAAGCTCAGTCAAGTTAATTTGGCCATTTCTGCAAGCTGTACAACTGTTGGAGCT
TGTTAATTTATCTTCTTCAGTGAAGAACCTGCAAGACTGTTAAATTTATGGGTAACATAAATGTTTATTCATAGGCAGACGGAAATGTTATAGATTGACTTAGGGATGCT
TGTAGGAAATAGGCTGGAAATGGCGAATGCTGTGAAAAAGATGATATTAGTTTTGAATGACTGTACTATTTCCCCAGAAAGAAACAGGTTACAGGCTTTAAGTTTA
TATGGGAAATGCAAAATTTGTTTCGAGCATTAAATTACTCATTTTATATGCTGGTCACAAGCTAAAAGAACTAAAAGGACCCACTAACGCATGACCCCAA
(SEQ ID NO: 118)

Exon: 6383..6317
Exon: 5566..5364
Exon: 3916..3768
Exon: 3678..3557
Exon: 3487..3261
Exon: 3188..2997
Exon: 1663..1001
Start ATG: 6383 (Reverse strand: CAT)

Transcript NO. : CT29768
ATGGCATTTAATCGAGCAGGATGCCCGCTGGAAATCCTGCCCGGAGCCGAGGAGGAAGCAGAATTGAGCGTCTATACGCGGCTCCCCTGAGATTGTGGCCCTGTTGTCCA
TTTTTCATGTGGGGGAATCAGTATGCTGGCGCGTCATTGGCAACACTTTGGTCTATCTGGGTGGTGGCCACGACACAGGCAAAATGCGGACCGTGACAAATATGTATATCGCTAATTT
GGCTTTTGGCGGATGTGATTATGGCCCTCTCTGCATACCAATTTCAAGTCCAGGCTGCCCTGCTGCAGAGTTGTGAACCTGCCGTGGTTCATGTGCAGCTTTGCGCCCTTCTGCT
CAGGCCCTGAGTGTAATGTCTCGGATTCTCGGATCTCAGCTTACGCCCATTCGAATCGATCGGATACGGCCATCATTAATCACTTAGGCGACCTCCCAACAGTTCTGATCGAAGT
TCATAAATTTGGTGGAAATTTGGATGCTGGCCCTGCTATTTCGCGGTGCGCTTTTGGCATTTGCCCTTTCGTGTGGGAGGAGTTGACCGAAAGATTTCGCGAGAACAATGAGACCTACAA
TGTGACCGGCGCAATTCGTCATGAACAAGAACCCTACCGATGATCAATTCGAATCCTTTTGCATACCGTGGTTTTTGTGAGATATCTGGTTCATTTCTGTGTGATCAGCTTT
GTCTACATCCAGATGGCGGCTACGATTGTGGGGACACAGTGTCTCTGGTAAACGACAGGATTCACGGGACATAACGCTGTGGAAAAACAAGAAGGTCAATCAAAATGCTGA
TTATCGTGGTCATTATCTTTGGACTCTGCTGGCTGCCACTGCAGCTCTATAATATTCGTATGTCCAGATACCGGAAATCAACGACTACCCTTCAATAGCATCGTCTGGTT
TTGCTGCGATGTGGCTGGGATGAGCAATAGCTGCTACAACTCTTTATTTGATGCGCATCTACAATGAAAAATTTAAGCGGGAAATCAACAAAGCGAATTTGCGCGCTGTTTCTGC
AAGTTCAAGACGACGATGCCGACGCGCAACGAAGAGGCTTTTCGATGCACACCGCGCAGCTCCATAAGGTCAACCTGACGCCAACTCTTCGATGCGAAATCCGAGATTAATCTCT
TTGGTCCGCGCGTGGTGGTGTCAACAAATGGGAAGCCGGGCTTGCAATGCGCGGGGTGCATGGAATCCGGTGCTACAGCGGCATTTACAACGGAAGTATGTGGGCGAGAACA
CAATGTCAATGGCCAAACATCATCAGATCAAAAGCTGGTTTACCTTTGCGGCCACTCCGGGTGTTTCGCGACCAGGTGTGGCGTTGCAATGCCGCGCTGGCGCGAACAAC
TTAAACCTCTGCATCTCGAAGCTAATCGAATGCGAGGACAGCTGGCACTGAGGACTGCCATCAACACGCGCCCGCAGGAGGATTTGCAATCCCGGGCGGAGTCCAGT
TGGCCCTGCTAAGCAGGGAGAGCTTCAGCTGCATTTGCGAACAGGAATTTGGCAGCCAAACCGAATGCGATGGCACCTGCATACTCAGCGAGGTGTGCGGAGTCCACCTGCC
CGGCTCGCAGCGCAAGGACAAGGATGCGGCAAGTCTTTGGGCAACCACTTTAA
(SEQ ID NO: 119)

Start ATG: 1 (Reverse strand: CAT)

MDLIEQESRLEFLPGAEEEEFERLYAAPAEIVALLSI FYGGISIVAV IGNTLVIWVVATTRQMRTVNTVNIYANLAFADVII GLFCIPFQQAALLQSWNLPMFMCSCFCFV
YALQSVNVSFTLTIAIDRRAHINPLRAREPTKFVSKFIIGIIMALLFVAPFAIRFVEELTERFNNENYNTVRPFMCNNLSDDQFSRYTLTVFVQLVPPCFISF
VQIQMAVRLVTRAPGNAQDSRDITLKNKKKVIIMLTIVII FGLCWLPLOLYNLI VVTPEINDYHFISIVWFCDDNNKNSCYNPFIYGIYNEKFKEFENKRFACFC
KFKTSMDAHERTFSMHTRASSIRSTYANSSMRI RSNLFGPARGVNNKGKGLHMERVHVGSGANGSIYNGSSGQNNNVNGQHQHQSUVTVFAATPGVSA PGVGVAMP PWRNN
FKPLHPNVICEDDVALMELPSTTPPEELASGAGVQLALLSRESSCICEQFGSQTECDGTCILSEVSRVHLPGSQAKDKDAGKSLWQPL*
(SEQ ID NO: 120)

Name: Tachykinin receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384568

AGTCGGAATTTTCGAAATTTGAAATTCGACCCCTTCGCCGCAAAAACTGATTGATTTTCGAAATTTCCCTCTGCGCTTTCTTGTTTGCTAAGTTTGAAAGTTATTGCTTTAACT
CATGCTGAAGAGAAATATCAGTTGCAGTCCGCTTTTAACTCTTTATTCGAAATTTATCTTAGTTGTGCAAAAAGAGTCTATGTCGTTAGTTGATGATATCAGTCGAT
ATCTCTTTAATTGGCCAAATACTGCTGAATATGTGGCAGCACAAATATAGTTACGAAAGTAATCGGCTTAAGAGCCACACTGCGATGTGCATTTTAGTGCATTAAGAGGACT
TGCAATGCACATCTGCTCTAACTCAGATTACGACGACAGCCATGGCTTTAAGCGCAAGTATTTAAAACTGGAATTTGAGCGCTGATTCGAGCTTAAATAGGAAAAAC
TGTTGGCTTTAAAGCAGCTGTAGTGTGCAAAATACCAATCTCGAGCATGGCTTCAAACTCAGCTCATTTCTGCTTCACATGACATGACATGACATGCGAATGGTATCAAAGTGCACACT
TGTAGACACAGAAGCCTCGGGTTTGCTTAAATTTTATGGCCTCAACACTTACCAGGTGCACAGCTTTCTTGCCAGCTATCTGATGTTAGTGCAGGGAATGACAGCT
GGAAGCTCGGATATCAAGAAAAATGAGAAATGAATGGAAAAATGGGTACCACCTCTTTGTTGTCAGACAAATTTCTCCTTTTAAATGAAGGCAAAAAGGATATATATTTTAA
ATAAGTTGGCTTCTGACAGACGATTCATACGCGAAAAATAAAAATTTAGTATTAATACCAATCGACTCGCTGGGAAGTATTGTCCTATTTTATTAGAAATTTTAAATTTTAA
TTTAGGGACCTTTAGTATGCTGCACATCAATCAATGACATAAAATATACGTTATTAATACATTTGGCTCTGGACACTTTGTGCTCGCGGGAATACGCAATTTAGATATATCTACTGAT
GTGATCTGCTGTATGATTTGGTGGATGAGTTGGCATCGACTGGAGCTCAGTGTGCGCTTTTGGACATCCAGCTCCATCTTGGTGGGTAATTCAGGACATCCAGCCACCCACGCG
ACGGTCTGTGAGTCCAGAGGACAAGTGAATGAACATCACAATCCTGCATAGTGGCGCTGATGCAGAAAGGATATGAAAAAGCCACACTCGCTGCATGAAAGCGAAT

56/89

Exon: 3391..2257
Exon: 2190..1001
Start ATG: 3391 (Reverse strand: CAT)

ATGTGCTTGAGAAACGGAACGTTTCAGTACTGTGATGGAGAGGAACGTAATTCATCGGACGAACCTTATGCAACACACATCTGGTTTACTGTACGTCCATGTTCTGCGAGCCGG
AGGAGTTTTCAGCAGCATACATACACGCGTACACACAGCATCAACACCTCCACAGCAACATGGGAACCGGCTAGAACTGGGCAACGCTGCCACCTCGACAGTGTGTTGCTCTTC
CGCAACCGGAATTCGCGGTCCAGCAGGAGTGACAGATGAAGGACCAACCGGACCAACTGGGAGTCCACCGAACCATGGAGTCCCGTGGTGTGTTGAGGCGATTACAGGAGCAC
ACGATCAGCGGTGACTTGAATAGCCTGCATGACGACGTTTCTC GAGGGCAGGAGGCGCACCAATGACACCCAGGGGCGTCGTGAGATGACTGGCATAATGAGGAACATGTTCC
GACGACGAGGTTGGGAATCTTCTACCCGCCGATGTTCCATCAGTCGCGGACAAATGTTCCGGCCGCTGATGACGACGAGGACAGGATGCAACGGTCAAGTGTGGACTTGGTCAGTGT
CTCAAGGAGATAAATGCTCTCGCAGATAAGTGTTCTAGATTATCAGCTCAGCTCAACGCTCAAAATCTTCCCTGCTACAGCGATTGCCAGACTACATGAGTGTCTCTCCGAG
CAACTGGTTTCCCAAGGACAGATGCGGCAAGGTTGGTGCCCAAGCCGACAGTGATGAAGCAGAAACGGCCACTACTGGAGTGGAAACGTACAACCTTCTCAGATATCTGGTGTC
AAGCTCTGATTACCGGCAATATCAGCGTGTCTCTGCCCAACCCCGAATGCGATCGCATAACTGGATTAGCTATATTCTCTGCCGAGGGGATCAACGAAGACTCTCCGCCAG
TGGCTTCTGTCACCGCTTATTCTGATTCTCCGAAGACTTGGCTAAGGTCAAGGAGAACTCAATCTGGAGCAGCTGCATTTTTCGGCGAGAACTCTGGCGGCAAGTGA
ACGAGGGGAGCAACCTATCTCATCTTTAAAGTCTATGCCACGATGCTCTGTCTGTGAAACCTCATTTGCAAGCAGCTGTAAGCGCCAGCAAGAAATGTAATCTTCATTTCAA
TACAGGACTGAGAAGCAATTAATCTCTCTCTGCCACCTGCCCTTTCTCTCTGCGTAATGAAAAATCTTCGCATTCGGATCAAAAGGCCCTTTAGCATGGGAGTGGATGCGGATAT
TTGGAACATGAAAACCTGGTCAACTGAGGAGTAAGCACTGAAAGCAGTTCGGACCTCTTGAAGATGCCATGATCGATGCCACACTAACCACCTCACACAGTTTGCATTC
CTGGTGGGTGGAAAGCTATCGAGCCAAACGACCTTGGCGAGGAGATCTTAATTAATCTCGGATTAATGAGAAAGTCCTGGACATTAATCTCGATAGTGGGCTCGAGCTTTCCCTTT
TGGGTATACCTGGGAATATCTCTCACTGCCGCACTCTTCAAAGAGCTGGCGCAGTCAGGCGCTCCACGAAGGTCTTACTGCACCTTTTGCTTGGCTATGTGTCTCAGATGATGCT
CTCTGCTGTTTCTCAATACAGATGATCTATCCGAAGCGCTTGTGTGAATGAAACACAGTTCGCTGTGTGGCTCTGGGCGCTGCCATCGAGTATCCATCTGTTCTCTTC
AGTTGGTGCTAATACGCGTTCTGCAATGTTCCAGCTTACGTACAGTATTCGGAATCGAGAGACCACTCGCTATATCTGTAAGCGGCCATTGTGGCCTGGCTACTGCG
CCTTGGTCCCCAGCTGCTGTTGGCCCTTATAGATCCGATTCGTAACGTCCCTCGGCTGCCCAACTTTCCACGGACACTGGCATTCGATATCCATCCGGGCTATGGTCTCAT
TTTTGGGGTGGTGCTCCCTGTACCTTGATTACTGTGTGCAACTTGGTGATATTCCTGTGATGTTTCTACAGCATCTCGCACTCTTTGAGCCAGAGACCTCAATAAAACGAA
AAGAAAAATGGTGGTTAAGCAAGATCCGACTCTCCATAATGCTCTCTTTTGTGGGTCTTACTGGATCTTTGGAAATATGCTGCTTATGCGGTCAGGCGAGGTGGGCTTTTCAAT
ACCTCTTCTGATCACGGCCACTATGACGGGATTTGTGATGTTCAATTACTTGTGCTCTTGGACTCCACGAACCGTGGGCTGGGTGGGTCTGATTTGCCCCACCAAGAT
GGAGCTGGATGTCCAAAGCGCACCACTGAGCTCCAGTCGATGACCACCTCATCCACCAACTACAACAGCAGATCACATCAGTAG
(SEQ ID NO: 122)

MCLRNGTFQYCDGEERNSSDELMTQTHLVCTSMFCEPEEFQHDYITRNHDQTPHQNKWKRRATIRATHDVCILRNGMEPVTRECRVKDNRANWESTEHWDDPVVCLRRFREH
TISGDLNLSHDDVLBEGRRMTDQGRREMTGMRNMFQRGGNLLPADVHMVTGFMGALMQQDKDATVSVLDVSVCKEIMSCDSKVLRLSQAQLNATSLLSQFESYMDALPE
QLVPKDRCGKVAUPTDEAEATATGVETYNFSDIGVQALITGNISVFANPECDRTGLAIFSAFGDQRKTSASGFWYRIFRESDLAKVKEESNLEATLAPENLWRQVKV
SRGATYILFKVVAHADLAFVETSLKRPKRSKVISIISIGLRSNYLSLPLPFLLRNENLRNLSQDSKAFSIGSGCGWYNYETWSTEGVSTESSNLLDKDAIECHTNHLTQFAF
LVGGSYRANLDGEEILITPINEKVLDDIISIGVCSLLGLIGLITAAFLKSKWSQATKVLHLCLAMCLQMMLFVFLNTDGEALVNGNNTVRVCALGAAMQYSILVLF
SWMLLIAFLQFQRYVTVIGIERPPPYILKAAITVAWLLPLVPTLLVALIDPSYVPSAAQLSTDGTGICYPSGYGLIFGVVLPVTLITVCNLVIFVYVFSYSHSLSQSIHKN
KMMVQKQIRLSIMLFEFLGLTWTGIFAFMQAGVAFSYLFCITATMQGFMVFIYFVLLDSTNRRLVWVGLICPTKMLMDVQKRTEQLQSMTSTSTNYNSRSHQ*
(SEQ ID NO: 123)

Celera Sequence No. : 142000013384531

57/89

Exon: 1001..1412
Exon: 2072..2174

58/89

Transcript No. : CT37292

Start ATG: 1

Name: Octopamine receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384349

Exon: 1001..1636

59/89

Exon: 1741..2208
Exon: 2283..2681
Start ATG: 1001

Transcript No. : CT37715

[illegible]

Start ATG: 1

MLNLILAIIIVFVISSQSEAVIPGCDYFDTVDISHIPKLNDSYAYEELIIPAHLTGLTYFRQLADGSGQEFVKSHLRACICKLKPCIRFCCPRNKMMPNSRCSGLTENLKRI
NPYLIKITLEDTGIGKYLLTDMIVLRYEFRYCEKVVSVQEDQYKLYEVKISIFLKYLSII*
(SEQ ID NO: 129)

Name: mth-like 6
Classification: G protein linked receptor

Celera Sequence No. : 142000013384207

AGTATATACGAGTATATCCTGTACGACTATATATCAAACCGAAAGTCGAAGTGAAGTGTCTCTTAAGCGGGACTTTTCATCGAGGTCGGTGCCCTCATATTGTAATTCATT
TGTGGGGCTCATTTGATTCATTGTGTCTGTGTGTGTGGCGTGTACTCTGTAAAGTTGGATGCAATTCCTGTGCGCGAGTGGTGTGGTTTGTGTGCATGGCCCCATCA
ATGGGATGCTCCATTCATGATCGTTAGTGGCGGCGAGCCAACTAACTTGTGAACATTCGTATGTAGTGTCTGTGGTGAGGCGGGCCCACTCGAGGCCCGCTAGCTCATGTAT
CCAATTGAACTGTGATTAGCTAACTAACTCTAGGTACCGACACCTCGTGTGCGCGTGGAAACAATCCGGCCTCTAAGGATATGAATTTCTTTACACTCGCTTACCTAATCGT
ACTGTGGGAATCGGAAGGAAGGTTCTGTGTGCGCGTGGGTTTGGTTTGGGTGGGAGGAAAGAACTTCTGTCTCGTGATATGTGATAGTTGTGGGACGCCCCACTTTGAAG
GACACTACTCTATTCTGTGAAGCGAATCGTATCCTTGGCGTTTTGGCTTACGCTTACTTTGGAATCAATCCGCTGTGCTGTGTGTGTATTATATCGCTACTTGGGTT
TGCTTAATGTAAATTTGTTAAAAATTTGTACCTAGCTTACAGCTTGCAATCCACCGTTTCATGCTCCACATAGCATATGTATGTATGTAGTTTATGTGGTATGTACAGATCTATT
TACATAAGTAGCTAGTAGAAGCTTTAAAGCTATTTGGGAACCGCAACGACGAAACGAAACGAAACGAACTTGGAGCTTCTGCTGCAACGATGCGCTCCATTTGGT
ACACTTTGGCATCTGCTTCCGGGGCTTTAATCCATTCCACAAGGACCATCACTAATGCTTATGCTATAACTATTGCTGGTACGCTGCTCGCGCTGTGAATACCCCCA
TGTGCATGATCTTCTTGAAGGCCCTTGCGAACCTCGGATTTGAATATTTGTAGATAAACCGGTTTCAAAAGCTGTTGATGTAGCCAGCAGCCAGTGTTCATCATGTAGGCCGT
GAGCCGAGGTCCGGACGCTTTCTTGAACCTTGGCGACATGGCGTCCATGTGTGCAAGTGAAGAAGCCGACGAGCAAGAACCACTGTTTGAAGGTTTGAAGGT
TATATATATAGTGGCATATGATTAGTAATGTGGCAGGCGCAAGGACGTGTCCAAACCGCTGGGGAGTGCGCTCAACTGAACTTACCCAAAACAATGGCCAATGT
GGTGCCCTTGGCTGCATTTTGTGCCGACGATTTTCCGGTTTCTTTTTCGAGCGCTTGTGCACTTGTATATCGTGAATCTGTGCGCTACGCGGAAGGATAATAGGCCATCGT
GTGGTATATATTGCTTTTCCACAGGCAAAAAAGGCAAAATGATGAAGAGGAGGTAGGAAGCGGAGAATGAAATCGGGTGGCATAGCATCGACTGGCCAAACAGAGCCCAAT
TAACCTAATAACTGTAATTAGGGGCAATGGACGGCGTTACTACCTTGACCGTCCGAGTCTTCTTATATCTTGGCGAGGACGTTTTCGATGTAGTCGATAGCGCTCGAGCTCG
TTGCGTGCCAAAGTCATGGCCCTCTGCACTCCGTACCGCACAAAGGACAATGGCTTCATGCGCAGCGCTGCGAACGGAAGTCGGTCGGAATTCGGAATGAGAACCTTAACCTCGCG
CATTAACGTACTACTGTGACTATTTTTTGGGCGCTTCCCGGATTTCCCTACATTTCCAAACATCGCGCTTCTATTTTTGGGGTTTCGAGGATAAAGTGGCAGGTTTCTCGCTAAACA
CTACGTAAATAGGCTCTAGTAGAAGACCGTGAGCTCTTTTTGCTAAGCAACTTGAACCTCAAGCGATTGTCAAGGATGCCCAAGAAATCAATTTGATGATTTTCAATGCGCTGCG
TGATATGTAAACATGCGCACAATAATATGAAGAGTAGTGCGAAGTGGCAGAACATAATCTGCAAACTCAATTTAAATGTGCAATTTGGATTTCAATCTGTTTGAAGAAATGT
AATGTGCTTAAAAAAGAAAAAACAATGTAATAGTTGACCAACAATACCCGCATATTTAATTAAGCTTTTGGCATTTTGTTTACCAATTAATTTTGAAGCAAGAGTTGTGACA
TTTTTGACCGCGTCAAACTCTTGTCTGTTTTATTGTAAGCAAAAAAGGATCGGTGTGTATACGATTATTGTGTTTTGTTTATAAAATATGACAATTTGATGTGTGCAATTTA
AGCAATTTTAATATGCTACGTAACTCTAGTGTAAAGTAGATAAAAAAAGCAATTTGCCGAAAAAGAGCATGCTCATAGTGGAAAAAAGATTTATATATTCAAAAAGATATGAC
AAAAAATTTGAATGGGAATGCTTTATTTTGTGCGGCTGCAAAAAAATTTCAATTAACCTTTTAAACTGTGCGAAAGTTTCAATTTTACCTATTGTGTACAATCAAAAAGATGTT
GTTTCAATGATTTAGTACCAATGTGGTAGACATATTGGACGGTATTTTGTGTTTTTTTTTTTTTTTTTCTAAATATTAGCAATTTCTCGAATTTCCGATATAAATGGCCGAACCTTGA
TTTTTAAAAACAATTAAGTTGGATGAGGCATATGTTTTTATCTCGTGCAAGCACTTCTACTACGCCCCCACTCATGGGATGCTATTTCAAGGTGGCAGGCTTACGCAATCTGA
ATGCCAAATACCTGGAGTTATGACACCGACTGACCTTGAAGCGGGGTTGCGCGCGCTTCGCGAGCTGTGCGCGGTGGGCGCGCTGACTGCTGACGCTTGAATCGCTGAG
GGTGCGGCCACTGGGCGAGGCACTGTCCGCGCGGCAAGCCGCTGTCCGCGAGCGGGCTGTATGTGCGCAGCAGAGGCGGAGGCGGCGGCCACGCCTGCAAGGACATCTGCAACG
TTTGAAGCTGCAATAACCAAGCATATGATTTACCTCGGCTCGGCAGCACAGCTCGCGGCTGTGTGATGATTGGGCGCAACAGCTGTGTCGGAGTCGGTTATGCAATGTGG
AGGCAATGTGACCGTGGGTTATCCAAATTAATGCTTGGCGGACCATAATCAATCAATCAATCGGCGATAGACACAGCTGGGCGAAGCAGCAGAAATCCGGCT
TGATGAAGGACAAAAAGATGGAATAATCGGGTGCAACGTCATTAAGGCGAAGCGCAAGATTGAGTTTATAAAGGTGACGCACTTAATTGAATTTATGAGAAATTTTCTCT
TGCCATTGGCTGTGATTTCTTTTGGCGGTTAAAGTTTCGCGGCTGTTTGTTCGATCCATTACCGCAATCGGTTGGGTCGGTTCGAGCTCGAATCCGATCCCTTTGCT
GCATTCCAGCTCTGTTGATTTAGCTTTGCTTGTCTTATTTGATTTAGTACAGGCGCATCAGGATGCTAATTTGGCTGGCTTAACGCTTGCACCTTATGTTGGCGTTTTCGCG
CAGGAATTTGGGAATAAGTTTCAATGCTCGGTTGGGCGGGGACATCCGATTGCAAGGATGATGGGCAAGGATCTTTTCGCGAGGACCTGATTGAAAGTGCAGACCGCTTTAT
GCGGCAATTTGAGTGGCGGCTGTGTCGGGAGGACGCTTGCATATTAGTTAGCAATTTTCCATGGGATGACCGCTTGGGGGTCGATTGCTTCTGCTGCTGTGCTGGTGGAA
GTGAGTGCACTGCAGACCACTCAGCAGAAACAGGCTATTCTATTAGCTGTTTGGGCGAGTGCAGAGGCTGTGAGGTTGTCAAGTTGCAAGTTTGAAGTTGCAAGCTGCAAT
AGGATATGGACACGGAAACGAACTCTGAGCTCCGAACCTTTTGCAGGGCCATCGGAGATAGCCCCAGCCAAACATTCGTTCTCCAGGCGATGGAGTTTTCGCGAAATTC
GCGGAAAGTTTTCGCGTTTTCATACCCCTGTCAGGTGAAGTGGTTTGAAGTTGCAATATATGCAAACTCCGACTCCATTTGGCAAGCTGTATATTCGTGGAAGTA
ACATTAGCTGCCCAAAAGAGGCTCTACACGCGCAAAAAAATTTTGAACCTGATATTTTCTTTTGTATATTTCAATTCATGATTGATGATGATGCTTCTCAGGATTT
CTACACAGCACATCAAAGCCCTAAATGGTATCGTAAAAAGTACTACTTTCGCGCTCTCTCGACAACCGGTGCCAGCATGAACCTCGGTGGACTTGTGTTCCAGATGAC
GTGGCACTGCTGATGCTCGGCGAGGCTGGGCTCTCGTCTCTCTCGTTGGAGCGGCTGGCGGTGTGTTGTGGCGGCTCTGTCGGGAGCAATCTTACGCTGGCGGAGCTG
CTGTCTCAGGCGCGTCTCCGCGAGGCGCGGCTGTGGCAATGTTTCTCATAGCTGCTGCGCGCTGAGTTCCGATAGTGTGGGCTGTGCGGAGGCGCGCTGCTCTCGCGCT
GACTCTCTCAGCGCTCTGTGGTGGGTGGGTACGAGATGTAATAATACAGAAAAACGATATACCAAAACATGGGACACAGGATACAGACTACGACAGCAGGACACGGACTACGGGCT
TGTGGGACCGGATCGAGGATCGAGGATCGAGGATCGAGTGAAGGCGGATTTAGGCATCGCATCTGCGCTTGATCTACAGGATTTCTGCCCCAGTACGACGCAAGAT
TGCAACACAGAAAAAGAAAAACAGGAAGAAAGAACAGCAATCCCAAGCGAGCATTTGAGATTGAGATTGATGATTCGCGCGCAAGTAATTCATTCCACTCGCATCGGAGAAATCCGCG

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Exon: 2186..1708
Exon: 1637..1001
Start ATG: 2186 (Reverse strand: CAT)

Transcript No. : CT38338
ATGGCATTTACCACGAGCTCATGCCCGCTTGGGATTATCGAAGCTCCCCGGAAGCGTTGGGTGTCTGCCGTTCCTGTGGCAGCACCAGAACGCCACGGAAACGCCAACGG
AAATACACCCCTCCAGGCGACCACTTTCGGTGCCGGCCATCTGCTGTGGCTAGCCATCAATGCCCTTCGTGTTTGCTCTATCCTGGGCGGGCAATTCTGACCATTGTGGCGGT
GCGCACCTGCCGCCACCTGCGATCGGTCACTTCAATCTGTTTCATCTTCTGCTGCGCCGCTCCGACCTCTGTGTGGGAGTGGCACTGCCCTCATCTGTTATTCATGTTTACATG
GGCTCGGCATTGGGCGCCATGAGAGGTCCTGCTGCTGCTGCTTCCTCTGCTCATCTCGCCCTCTGCGTGTCTGATCTGATCTCGATTGCGGTGGATCGGGTATG
TAGCAGTCGTCTATGCTCTGCATCTAGAAATACATGACCCGTCGAGTGGCATATAGCATATCATATTAATTGTTGTTGCTTGGGTGCATTGGTGGCTCTGCTGCCAGTGT
CTGGAATCGATGCCCGGATGCACAGGCTGCGCAATTGCACGAAGTTCTCGGCCCTGGTTACATTCCGGGAGTGATAACGCCGGGTTTGTGATCATCTCGATCTGTATGTTCT
CTGCTCTACTGGCGCATCTGCGCGAGGACCTCAAGCAGCGCGCTGCGATTGCGACAGTCCGTTGGTCTACAAACAGCAGGACGAGGCCACCACTTGCAGAACTCGCTGCTCAT
CAGACTGGAAGAGCGCTCGGATGTTGGTCTTCATCTACGGCTGCTTCACGCTCTGTGGCTTACGCTATCTTGTTGTGGCCATTGCCACGCTATTTAGCATCTGCCAGAGCAG
CTCGATGATCTACAAGACGACGTTCTCTCGGCCATTGCCAACTCTGCTCTCAATCCGATTATCTACTCGTGGAAAGAACTCTGGCTTTTCGGCGAGCCTTTGTCCTCAACATTA
TGCTGCCGAAACGGCGAGGCACTGCGAGGATCAATTGCCGGCGGACAGCAAGCACCAGCTGAGGCCACATCCTCATCTCAGGAGATCAACCCGAAGGCCACTCTAGTGA
(SEQ ID NO: 134)

Start ATG: 1 (Reverse strand: CAT)

MAFTSSSSPAWDYRSSPEALGVVPFLWQHQNATETPTTEITLQATSGAGHLLWLAINAFVLVLILGGNILTIVAVTRCRHLRSVISNLFILSLAVSDFCVGLALPYHLVLFYM
GSDIGAMRGPCLLRFFLLICACCVSMLTLISIAVDRIYAVVYALHYRRYMTTRRVAYSIIIFNWCGLGALVALLPVEWNRWPAQACEFEDEVLAGPYIAGVITPGFVIIWICMF
LVYWRIMREASKQALRLRQSVVYNTDEATTMRNLLLHPDWKSVQIVVFIMGCTLCWLPYFCVAIAQLFSICQSSSMIYKTTFLAIAANSALNPPIIYSWKNSGFRRAVQTL
CCRARQCEDQLPADSKHRMEATSSSQEIKPKATQ*
(SEQ ID NO: 135)

Name: Adrenergic receptor- like
Classification: G protein linked receptor

[illegible]

CCAAATGCTCATTGCACCTCATTGCTCATTTAATGCGGCACAAATTGACGCGGGAATGGGAAAAATACATTGTGTCGCCATCTCCCTGGTGACTCGAAGCAGGATATCGATAAAT
TTCCCAATGCGCTCGCAGACAACTTAGTGCTCTGGGTTTGTTTATCGCACATTCGAATGGAAC TGGGGCTCAAGGATAAAGGATAATGGATGTGGCGGACTGAGGGCGGAGATCC
CATAGACGGTTGCGCCAACTGACAGAACTGTGTTTAAATGTGTGCGCAAAGGACAAATGGGTGCAGATACGCTGGGAATTTGGCTGTGCCCTAAAAAAGTGAATAATTA
AAAGTTTAATGACAGACCAATGGGCAACACCAGCCCGGAAAAACGCATTCCTCAACAGCTGTAATGAAAAATACAGAGAGTTTTCATTCAAATAAAATATAAGAGTATAAAG
GAAATGAGTCGTTGTAGCGCGCCAAAATAAATAAAGAAGTATTATTATAAATAATACCATCAGTATATTTCTCAGTCATAATGTTGTATGGCTTTAAAGAAAGTTTGAAGAAGT
AAAGTAGTATAATAAATAATTTATTAGACAGAGGTAAATTAAGTAAGTAGAGAAGGATTAAACGAAAACAATTAACCTTATGAATTAATGTGCTGAAAGAGTAATCAAA
CTAGATTTCAGTTTGGCCAGATGTCGCAATCAATGCAATTGGACCAAAGGAGAAACGGAAAAATGGGAAAAATGGACAAATGAACTTTGTCTTTTGTGCGCAGGACTTTTCTGTG
CCTCTTTTGGCCCCATCAACAGCTGCCCTCACATGCCCTTGGCCCCGAATGCAAAATGAGGCCATAGACAAACTTGCAAAATTTGGCCATAAAAAATTTGCCCAAAATTTGCCGCCCA
AAGTAATTCGATTCGCATATTTGTCTTTGCATTGAAACAGATCCTGAAGTCGCCCGCATCAGCTGCACATGAATCTGTTCGCCCTGTTCTGCTGCCAATAACTCTGTGTGG
CTGGCTGTGGTACCTGTGTGGTCATGCGCAATTCGGAGCTACTGCATCAGAGTTCGGTAAGTGTGCACACCTGGCGGAATGCGTAATGGTGAATCAACCTCGGGCTCCGATTCC
ATATTCCATCACTCCGTGCGCAACCCAGATGGCGTTCGCTTGCCCTGACCATACAGCTACACTACTCTCTCTGTCACATTTACTCTCGATGCTCTGCGGAGGATTCATCTGC
CACACCTCTCTGTGCGCGCAATTCATTTCCGAGAAGAGCGTGTGTAATGGCTCATCGCATTCGGCTGGGGCTCCCCGCCATCGTCATATTCGCTATAGCATGGCTCGCG
GTCTGGCGGCCACGCCCGAGGACAATCGTCAGTAAGTGAATAATGAATGGGTACAGCTTACACGAAAGAAATGAGGAATGCCAATAGTTATCAAAAGTGACCGACAACAG
ACAAGGTGTGAACCAATATTGTGTGCTAGAGAATTCTCTAGTTTAAATGAATAATCCCAACAGACTTGAGTATCAGAGTTGATATCAAGATATTTCTGTGTAGCTAAGATATTCTAG
TCTGTCGAGCTTGTGACCAATGCTAATCAAAATCAATCGCAATTCACCACTGACTCATCTCGCATTCCTTCCGCAACTTCGACGTGTGGATGAACCAACCACTACCA
AAACATTTCTTATGGTGGCTGTGTATCTCCATGCTTCTGAACTCTCTGTCTCTGTGCACATCTGTGCGAGTGGTCTTTTGAACATGAATGCCCGCGGCAGTATACAGGGC
AGCTGCGGCTCCATCGCGCAAGGTTTGGCAAGCATTTTCGGTAAGTTTTCGGCTCCAAAGCAAAAAAAGAAAGACTGTTGGGGGGGAGATTGAAACACTTATAAGCGAA
GCAGAGAACTCTCAAC TGGCCCTTGGCGGAATGCGTGAATATATTTTGTTCCTCTGTATTTAGACATCTTTCCCATCGGCCATCAGGCTCTTTATGTTTATTTCGCTCTGC
ACTCTGGAAGTTGGGCTCTATTTACATTTTGAATACGCATACGCCCGGCTGACTTGCATGTGAATACACAGCTGAAACACGAAACGAAGGCATATAATTCACCCCTCTCAT
CCTGTTTATAGGCAACGCTGCTTTTGTTCTCTGTCTCGGCCCTGAGTACATCCTTACGCCCTTTCTGCTCGCACCAACTCCCTGGGAGAAATACATACGAATCATCTC
GGCGTTTACGGCCTCATTTCAAGTAAGTAATAATGTTTTCGTTTCGTTTCTGTCTCGTTCGTTTCTTTCGTTTTTTTTTTTTCGTTTTTTTTTTTTCGTTTTTTCTGTTTCGCGCTTTGTTTTC
CACGCTCTCTATAAATCACTTTACGATCCGAGCTGCGTAAAGAGTTATGATGTAGTTCGCCCTATGTTGGTGCATCCACCACTGGGAATGCAAAATGCGCAGCATTACTGGA
ATACCGAATAACCAATACAGGACTACGAGCTCATTAGATCGGGGCTTTTTCAAATCAATAATGAAAAATACAGCCATGTGAAATACCCCAATTTGGCATTTCCATTTGATC
TCAGTTTATTAATTTGCTGGCCGAGATTGGAGCGAAGTTCGCCGTTGGAAAAATGTAGCAAGTTGAGCATGAAATGCACCTCTCGCACACAGGAACCTCGTAAAGCAACTCG
AGCTGTGCCCTCGGCCAAGCTCAGTCTCGATAGACAGCATTTACATGATGTGAACACACATACATATTTTCTGGCGCAACTTAAAGACAACGCGGCAACTCATCCA
GCATGTTGCCCTCTCTGTTGTGAATACCAAAAGATCTTTCAGTTCATGTTTTCGATGTCAGATACAGATAAATAATTTTGCATATATTTTCTGTTTGTGTCGCAAGAAATGGTT
TTTCATTACAGACTGTTGATATAACAGTTGCCATGGAGGTAGAGGTGCCATAAAGTATGTTGCAGCTTATAGTGGCTTTCCTTTACAAAATTTCAAAAGTAATTTCTAAGGCA
CCCCTAGCTGTTTATCGCAAAATTTTATGTCGATCTAAACCCCACTCAATCGGTTCAAAAGGTAGCCCATGTAATTCATTAACAGGCCCTTAAACACAGGAAATCCCA
TTATCTTTCGAGTAGACATAAAATCATCTTGGGTGCACTAAGCTGCAGTTGAGCTCTGCACTATTTTCTATAACTTGGCAAACTGAACATTGTCACCTCGGCTGAGCCTT
TTTTCATCATTGCCAATCGATAGGTGAAATCTTTTATTCGACTGCTAAAGTTTAAAGCCAGTCGACGGGTACTTTTCTTTGCATTTTCTCTCTGTGCTATTTATTCACCTT
TTTTTATGTCTGCTGTTTTCGGTTGGCTTATTGTGTCGAAGA
(SEQ ID NO: 136)

Exon: 1001..1196
Exon: 1590..1727
Exon: 3775..3881
Exon: 3955..4046
Exon: 5156..5318
Exon: 5405..5631
Exon: 5910..6086
Exon: 6395..6659
Start ATG: 1031

[illegible]

Start ATG: 31

MSDQIGNPNATFSGSGSGSGTNVASIAESVAESGPDFDALRAACETRLNASQGLAGSGGPAEAGTHCAGTFDGLWLPDPTAVGTSAYELCPDFFITGDFPARYAHKECGLDG
EWFKHP LTNK TWSNYTTVCNLEDLNRWHTVNLISEVGYGTSLLAILLSLALGYFNNSIRIFVFLRNLASLKCARITLHMNLFASFAANNSLWLWVYLLVMPNSELLHQSPMR
CVALHITLHYFLLSNYSWMICEGYLHTVLVAAFISEKRLLVKWLAIFGWGSPAIVIFYSHARGGGTPEDNRRHCWMNQTHYQNILMPVVCISMFELNLLFLCNIYVRVVLKLL
NAPASIQGSCGSPRTVLQAFRATLLLVPLGLQYI LTPFRPAKHPWENTYIISAF TAS PQVSKMFSFRCSRFVFFSFFFFFAFFFPFVLHALIKSLYDSELAKKL*
(SEQ ID NO: 138)

Name: Calcitonin receptor-like
Classification: G protein linked receptor

ATTCGCTCGGGTATTTTCGCATATGTTTGTGGCTAAGATATCATCAATTCAACACAATGATCAACAATTTCAACGCACAACTCGACTTGGGACCTCGGATGCTTC
ATCGGGGGCGCTCGGGAACATGAATTTGACCCATCAAGGGAACACTGACCAAAATGCAACGCTCTGTGGTAGCGCATCAAAATCGAAGCGTTGAGGGGGCGGCCAAAAGGAAA
ATGAATCATCTTCAAGGATTTGAACACAAGTCAAGAACTGCACAACTAGCCATCATTTTCAGTATCATAGTTGTATGTTGTAAGCGCTAAGAACTACTGGAATATGCTCTTTGGGG
TTATTTGCTTTTTCGGATTTTAAACATATTTGCTGTGTTTTATATTCGTTTAGTGGGCTCTGGTTTTGCTTTTCTTAAGCGCAATTCGCAAAACGGCTGCGAAGTGAAGATTTT

[illegible]

Exon: 3185..2737
Exon: 2223..1965
Exon: 1876..1001
Start ATG: 2869 (Reverse strand: CAT)

Transcript No. : CT41076
ACACACTACACATCGACACACTCACCTAAGCACACATAGACACTCACAGACACACCGACACACAAACCGAAGGCACAGGACAATTGCATGGGCGTCCGCCATGACCGGAA
GTAAAGTACCGTCACGTCACAGTCCTACTGGCCCTTGCATAAACAATCAGCCGCCAAGAGAGTGGCGAGAAAGCGGGTGAAAGCAGCAGCGAAAGCAGGTGAAGACGAGGAGGA
GCCAAGGAGGAGGGAACCGGACAAGGAGCTAAGGGAGCGTATACAACGACGACAGGAGCGCCCATTTCAACCAGACATGGCCAGGCTAAAAATGCAGGAATAGCTTACCT
ACAGGATAATCCAAAGTGGAGGCCCTCACCAGGCGGTCTCATATCGATCGATCTGGCGCTGCCATCGCTCTTCCAACCTCTCTCATTCGCCCACTATGCGCACTTCAAG
GGACCACAGAGAGTGTCAACTACTCTTGTGCTCGGCCATCGCCGATCTGCTCTGGCAGACTCTGCTGGTGGCCCTTCTCCGTGTATCCCGCTTTGACCGGAGAATGGA
TGTACGGCGACATCGTGTGTCGTTTACC GGCTATCTGGAGGCTCATCTGGGCGGATCATCGGTATACACCTTCATGTGGATATCAGTGGATCGCTATCTTCGGCGTGGGAA
GCCACTCGATATGAACCGTGCAGACGAAACGAGATGCCAATCTGGATGTGGTGTTCATTTGGATATCGGCGCCCTGCTGTGCTGTCCCAATCCTGGGCTACTCGATG
CCCATCGAAGAACACATGACGCACATTTGCATGCTGGACTGGGGGAATATGGCAGCATATAGCGCCACCTTGGCGATATTAGTCTTAGGTPCCAGCCTCATTTCATTCGTAC
ACAACTACCGGCTATCTTTGTATGATGCGGTAGATTAGTCCGGCGGCGGCATACAGATAAAGAAATATGCAACTGCTCTGGCTGAAAATTTATGCAACCTTAGTCATAT
GATGCTATCTGCATATAGCTCTTTGCTACTCTGGGTGCTCGGCTGCCATGGATTTCTGTGGCTGTACGAGGTGCTACGGGCGATGTTATCCAAATGACTCTTATCACTTCT
GCCGTCTGCTGGATCGGCATATTGAATTCGTTTTGAAAAATCTGATCATGACCAGTATGTCGCCCAATTCGGTATCGGTTTGAAGATTTTGCCTTAACCAATTTGTGTGA
AGCTAAGGCGCGTTTGCAGACAGAGCTAATCGGGTTGGACCCAGATGACAGTAGAGTTAGATTTTTCCTCTCCCTAAAAAATAACATAATTATCCGTAATAAATC
CCCCACCAAAACCATCCCTCGTAATCCCTTTGAGAAGACTACAAGTCTCGCAAGTAAATATTTGGTGTGAATTCGATTAGGCCATTTCCCTTAACCTCTCTATATCTCT
CTCTCTCAATATATATAAACTTACTTGCTGTTTGCCATATAGTTGCATATTTACGGGCGGTTTGGCGCTACGCGACATGCCAATGAAAGCACCCACCCACAGTCACAGC
CACAGAAAACCTGAAA
(SEQ ID NO: 140)

Start ATG: 317 (Reverse strand: CAT)

MQEMS YLQDNSKVEALTKAVLISILGVAVLNSLLIIATYANFKGPTEVINYYLLSLAIADLLCGLLVVPFSVYPALTGEWMYGDIVCRFTGYLEVTLWAVSVYTFMWISVD
RYLAVRKLPRYETVQTKTRCQCMVFWTWSAALCCCPILGYSMPIENNMTHTICMLDWCNMAAYSATLAILVLGPSLTSIVHNYGYIFVMRKRIRSGEPIDHKEYATALAEN
LSNPSHMMSFALVFAFVWSWLPWILLRLLEYVVTGDVIQSTLINFVAVWIGILNSFWKILIMTSMSPQFRIALRVFCLTICCKTKGRQLQAEILIGLOPDD*
(SEQ ID NO: 141)

Name: putative adrenergic receptor
Classification: G protein linked receptor

Celera Sequence No. : 142000013384685

TAAATATTCGCTATTGAACGAGGCATGCAATGCCATTGAGGTCCGGCTGTGGCTTTCAGTTTGCATTTGAACGAGGATTATATTCAGATTTAGCCATTAGCCAAATGACACACCTTTCGACGAGCTGTGACACGTCGATCTTTGATTTCGGCCAACTGAGTGATCCGGGTAGTAGTGTGACATGCCAACATTACTTATGCACTGAAGAAATAAATAAGGATTTACATGTGTAGTAATGTATAATTGGCGTCATGACATGCATGATAATGATCAACTTTCTCAAGCAATAATAATTTCTATGACATCAGCAGATTCATACGGACAGAGAAGACGAACATGGATTTACCATTTAAATATGTAATATAATATAATAAGTATCTAAGAAATTTTTCATCTGGATCTCTCAGCAATTTTAAACCATATCTCTCAATGCCAAATGTGTAAGTTCTTGCAAATAGATACAGACAGATCAATGGTCCGCAATTTGGCAAAATTAATTTGCCCTACAAACCGCAGGTGCTTTGAAATCAGAGTGCACATTTGGACAGGTTTGCTCA

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GCTCACTGGCAGCTCCACTTTGGCCATTCAAATTTGTTTATCGCTTTGACAGAGTTGCAGTAGCAATTTGCACCCGATGCCCAATGAAGAAACGGCATTGATAATAACA
AAGTAAATGATAAATAGATCGTGTTGACGCTCAATTTCTTTTGTATGATAGTATTGATTTATACATAAATGACAAATTTCCCTTTATTTTGTAAACATTATGCCATT
TCACTTTAAATGGTCTGTGAACACCAAGTTAGTTGTAAGACGACAGAAAATCTTCATTTCGATGACAGAAGTCCTTTAAATTCAAATAATATAATTTCTAACTCTAA
ATAAATAATTTCTAAGCAATGGTTTATGTATTTAACTCCTTAAATTTTGTTCACGTTTTTAACTAGCACATATACATCAAAAGAGCGGAAGTATAGTAGTCTACAATGATGCA
AGAGACCGGCATCAATAGTGGGACAGCCCATATGACACAGCGGGTCTCTTCAACGACACGGTTCTGAAAGCATATCATCTCAGCAGACCCGATATCGAGAAGTTTGTAAAG
CTATGGCAGGATGACCAATGAAGAACATGACCCCTCAGGTGGATGAGTGGCAGGGCTATTCGAGGGGAATATACTAGCTGGTGGCAGCTACACAGCAATCTCATGGAAT
ACGTTTCGCTGATGGTAAGTAATACTCTGTGATTACAACCTCCCTTGGGCACGAACCCGCAATGTTGGCCAAACAAAGGAATCGGTGACTAATGATGAGCAAGGGAAGCG
AAGCACTATACACAGACCTTCGCCAGACAGCACTGATGAATGTAGGGGACTCTTACCTTTGAACTCTTGCCAAAGTCGTCTTTTTCGAATTAAGATTTATAGCATGGT
GCTCAGCTTAACCATCAAAATCAGAAATGAATTTAGTGGCGGGAATCTCGCTCTTTGTTTATGTCCGGGTTCCCTTTTACGGGCTTTTTTGAAGAAAGATATAATTTATA
TTAGATGTAGTTAAGCCGTTGAGGGCTTAATGTGCTTTTTCTTATATAATTTTCAGTCTGTAAGTCATTTTGCCTTTGACGAGACGGCCTTTACCACAGGAATTTGGGT CAGAA
AATACCGAGAATTCATTTCCATCAAGGCTAGTCTATGCCCCATCTTTTCAAATAAATATACAAATCAAAATCAAGTTGAAAAATGTAACTTTATGGCTTTCCCTCTGTGCT
CAGAAAATAAGTTTACCTCTTAAATTTGAATAAAATCTGAAACAACTCGGTTACTGTAGTTCCATTTTTTTTTTTTTTTTGTGTTGTGTGAGCAATGTTAAGAAATGTAA
ATACGAGAGGTCACGACAGTTGAGTTGCTGCTCAGGAGCTGAATAAAATAAATTCATCTCACTTTTGGTGCCAAAGGGGAGTGCAATTCAGTTGAAAGTATGTGAGCGCTCAC
AGTTTCGATTTGACAAAGGACGCATTTATGTTGATTTCTTTCTTTTAAATCAAGCTAGATTTGCAATTTTCGGGACATAGCAACATCTTGAACATAATGTTGCTTGA
CCAGAAAGGAATGGCCAAAGACCCCAATAAATAATACTAATGTTGCTGGCGGTGGCCGATGTTTTGTGATGCGGAATATATACCCTATACATCTAGTATATATA
TATGGGGCCAGTAAGACATACATTTAGTAATAATTTAATAAATAGTCTTGGTTATAGCTTACCTTGTATTGTTATTTATTTATAGTTTATAGCTGATGATTATATTATAAAGGT
ATACTCTTGATCAGGATCTACACACGCTGTCTGCTCGGTTTCTAAGGGTATATTTTGTAAACAATTTGTTAAACAATTTCCCATATATCTTTCCCTTTGCACTTACTTTTA
GTGAGTAACGAGTATGTGTTAGTCGAGGCAATCGGCTATGACCTTCTCTGTAGGTTTCTTTCTTTTAAATTTTAAAGCCGTTTAAATTTTTTGAATTTATTAATATCT
TTGATAGGCTGTGTGAGGTTGAAAAAGATTGAGCTACACCTGGGCGGTTTGTCTGCTGGTTACATGTCACCTCACCAGCATCTCGCATACGATTTCCATCGGATTTAGCTGTC
ACGCTGGCGGTGTGGCGGTATGTGCGCTATCAGGTGAGATGTTCTTCCGCGACATATGATCATCTTCCGAATTTCCGCGTCACTCCATGTAGAGTTTGTCTCCGCTGCGC
ATTCATGATAATCCACATCCGCAATTCGAACAGACATCCGGCTAATCTGATTACCACATCATTGTGGCAATGTGGCTTCTCTCTCTTCCCGCGTCTTTTGTATTATAGCCAA
ATGTTACGCTCTATTATCCACAGAGCTCCGAAGCGGGGCTGTGCAAAATTTCTGCTAGCACATTCGGGAGGCGCATCTCTTCCGCTCTCATCCGTTGCTGCCGATCTCTGCC
TGCCACAGTCAATTTGTGTTCCAGGTGCGTGAGAGCTACGAGCTGCAAGGTCACTCGGAGGCATGTATCATGTTTTCGCAAGGACATCCGGTGTCTACAGGTGCGC
TTAGATAAGAAGTCGATGTCATTTATGCATCTTGCGAAGAACCAATTTGTAATATTTTCATCTGTGAAAATCTTTAAACAGATTTCAACTCTCGGATACATTTCCGTGCTCATCA
AATCTTCGCGTGGCGCATATTAATTTGTATCGATGCAAGTACTTTATGACGTTTGTGCGAGGATCTCGAGAGCGTCTTGAAGTCGAGGATTACAACATCCCGCCCAATA
TGCCATCCAGCTTAATCTGAATGAACCAAAATCCAGAAGTAAGTACGCACTGCTGTTAGTGCTTTAAGAAATCGCCCTCATTTGCAATTTGCGCTGTAGTTTCTCATGAT
CTTTGATTTTCTGTTTTGAATGCTCTCGGATTTTGTCAAATCTCTTCTAGGTTCCCTTTTGCCATCTTTTTATAAATTTGCTCTTTTGTGTTTGTATCTTAGGCCGCC
TCGCTGTGATCGCGCTGAACAGATCGGATACCTTTTGTGCTGTTGCGGTATGTTTGTGTTCTGATACGAGGATTTCCCGCAGGGATTACTTGGTCTGTGCTCGGAGTAGT
GAGAGTGCTCTTTCCGCCATCTGATCCCGCGTTTGGAGAGCTGATGATGCTGTGCACTGATCAACGCGCGCTCGGATCTGCTGTACGCTGTGATGCTGCAACAGT
TCCGGACCACTTCCGATCGCTGTTCAATGAAGCGGCACCTTTGGCAGCACCGAAATGACAGATGACCCGCGTAAACCAAGCATTTGCGCTAGTTGGCGCACATCTAAAGGAT
GATCATGGGCAGGAGCAACCACTAGTGAGGATGAAGTAGTGGAGGACATCCAAGATCTCCGCGCGGCATTTGAATTTCTAAAGCATAAACTTTATAAAGTT
TCGTTGACTAAACAAAGCTAAATATGAGAAAGTGTGAGTGTGGAATTTATGACGGGCTGACAGGTGATCAGTTGATTTCCCAATAGTTTACTTTTAGGTGATGCTCTA
AGGCGCTGCACTGGTGCATCGCTGTTTTGAAGATTTGATTCGTTGTGATCTGGCTGTTTCCGAAGTGCCATTTCAATTTCAAATAATTCATGATGTTTACCAGTTG
GTGTTTCAATTCGATTTAGTATAAAGTTTCTATAGGCTAAGTCGAGTGTACAAGCTGTACAAGTACGACATTTACTTTTCTAATAGTGCAGAAAATAACAGGAAT
CCATCATCAGCAAAACTGTAGCCATGCTAAGAGGACTGAAGAACTCACTGAGCCCCGAGTTGCTCTCTCTTCTAGCTGGAATCAGGAAGAAATTTCTATTTTGATTGTATGG
CGTTAACTGGTTTTCACAAAGTGCAATGCGATTTTAAATTTGGCTAATTTGAATAAATTTTGAAGTACTTTTAAAGTCGCTTATCTATTTATAAATAAATGTAACACT
TCAAAAATAAATATGTTTGTATGCTGAGTTGCATACAAATGAAGACTTTTATGCTTCCATTACCCGAAATGAACATCTTAATCTGACTGACACAGCAATCTACCTCCAGTGG
CAATTCGCTCCGCGAGAAGTTCCGTGGCCCCGATTGTGTCCAAATCAGACTCGCCCATATCCGCGGAATCCAGCAGGATGCCATTGCCCTGATTTCGAGATGCGCAGCAGGGAT
TTCAGCTTTGGTGGCTTGAATTTGTACTGCAAGATAATGAGATGATAAAATCTGCTCTCGAGGGAGTTTCGACCATTAATGTA

(SEQ ID NO: 142)

Exon: 1001..1246
Exon: 2080..2251
Exon: 2594..2720
Exon: 2936..3130
Exon: 3217..3399
Exon: 3579..3900
Start ATG: 1001

Transcript No. : CT32473

ATGATGCAAGAGACCGGCAATCAAAATGGGACAGACCCATATGCACCAGCGGGTTCCTTTCAACGACACGGTTCGAAGGACTATCATCTCACGAGCACCAGATATCGAGAAGT
TTGTAAGACTATGGCAGGAGTACCAGATGAAGAAATGACCCTCAGGTGGATGAGTGCACAGGGCTATTGCCAGGGGGAAATACAACTGGTTGCGAGCCTACAAACAGCAT
CAATGGATACGTTTCCGTCGATGATTTGCATTTTCGGGCAATAGCAAACTCTTGAACATAATGGTCTGCACGAAAGGAAATGGCCAGACGCCCAATAATATATCT
AAGTGGCTGCGCGTGGCCGATATGTTTGTGATGCTGGAAATATATACCCATACATCGTATCAGTATATATATATATGGGGCCAGGTGAAAAAGATTGAGCTACACCTGGCGCG
TTTGTCCTGCTGTTTCACATGCACCTCACCCAGATCCTGCATACGATTTCCATCGGATGACTGCTCACGCTGGCGGTTGTGGCGGATGTGGCTATCAGACATCCGAACGGGGG
CTGTGCCAATTTCTCGTAGCAGATCCCGGAGGCGATCTTCTGCCCCATCTCCTGCCCCTGCTGCTGCCCAGCTACTTTGTGTTCCAGGTGCGTGAGACGTAC
GAGCTGGCAAGGTCAACTCGAGGCCATGTATCATGTTTATTTCCGACAGGAACTCGGTGCTCATAGATTCAACTTCTGGATACATTCTGCTGCTCATCAAACTTCTGCGGT
GCGGCATATTAATTTGATTCAGTGCAGTACTATGCAGCTGTTTGTGCGAGGACATCAGACGCTGCTTGAAGTGTGAGGATTTACAACAATCCCGCCAATATGCCATCCAGCT
TAATCTGAATGAACAAATCCAAGAAGCCGCTCAGCTGTGATCGCCGTAAACGATCGGACTACCCCTTTTGCTGGTTGCGGTATGCTGTTTCTGATATCAGGAGTTCGCG
CAGGAGTACTTGGTCTGCTGTCCGGAGTGATGAGAAAGTGCTTCTGCGCACTGCTATCCGCGGTGGAGAGCTGATGGATCTGCTGGCAGTGATCAACGCGCGCTCG
GATTCGTCTGTACGGTCTGATGTCGAAACAGTTCGGGACCACCTTCGATGCTGTTTCATGAAGCGGCACCTTGGCAGCACCGAAATGACCAGATTGACCGCGTAAACAC
GACTTGCCTCTAG
(SEQ ID NO: 143)

Start ATG: 1

MMQETGNQMGGQTHMHQRPVFNDDTVLKDHYLSTSDIEKFKVLWQEYQMKMNTQVDEBCQGYCQGEIYNWLRAYNS IHGYVSLMICIFGTIANINIMVLTRKEMAKTPINNIL
KWLAVADMFMVLMLEYIPYTSYQIYMGPGEKDLSYTWAVCLLVHMHTQILHTISIGLTVTLAVWRYVAIRHPNGGCANFLLAHSREAILPFIILSPILCLPTYFVQVRETY
DVDRVNSAMHYHVYFDKDSVLRYRFEWTHSVLIKLLPCGILTVISAVLMHVLCASRRRLKLRDYNYPAKYATQLNLNETKSKKPPRCRRNRDRTLLLVAVLVFLITEFP
QGLGLLSGVMEKCFEACYPFPGELMDI.LALINAAGFVLYGLMSKQRTTFRSLFMKRHEGSTEMTRLTRVTTTCV*
(SEQ ID NO: 144)

Name: TRH receptor-like
Classification: G protein linked receptor

65/89

Celera Sequence No. : 142000013384808

GTAAATGATTACAAACAGCCGACATCTCGGGAACCATTAAGAGACATTTAAAGTCAAAAGTGGCCAACTGAAACCCGTTAAATCCGCTTGGCACCACCACTTGGCC
CGACTAAGGCAAAAAATAACATAATCTGTCAAGTCATGCAGACAGGGCGATTACATTATAATTTAAATGGCAATGGATTAGTCATCATCAGTACCTGCTTATCTGGTTT
TCCTGATTTTTAAATCAAACTGTGACTGTATCTTTACCTTCTCTTGTACCTCGTTATTATAAAATGCGAAACCCCTGCCAAAGGACACTATAAAACGAATTTTTTA
AAAGACGCCAAACAAATGGCACATTTTGTGTGAGAAGAAAATATGCCGATAAGAACGTTTACTTGTAAAGACGCTGCAGAAAGATTAGGAACGAAAACATATCGACAATGTTT
ACTTGTATAATCTGAGCTTTCTAAAACACAAAATAGTGTACTTTTACCATACGCTTAATTTTGAAATTAATCCGGGATATACAGTTGGGGATCCAGATTTTAAATTCGTGA
ACACAGCTCGCTGGAGTAGGGCAATTTCTGTATCGATCTGTTAGTTTTCGGACTTTTACTTTGGTTGTTGTAACAAATATCCGCAAGTTTCTCGCAAAGTGGGTTGCGTTGTGC
CGTGCCTCAGTAGAGCACTACACACTACCTTCGATTGAGTTCGGTGGGATTGGATTTCGTTACAGTCCAAAGTTTGGACCGTGTAGGAGAAGATAATGATCCGATCACTC
AATTAACCTTCATGCACCCAGTGACGTCTCATGTGCTTACTATTCCGTTGCTATATAGTCTATTTTGTAAATTTCTTACACCTTACATTTTCGATACGATTTCGATTTTCGTT
TGGAAATTTGCTGAAGGTAAACAGCATAAATTTGACGTTCAAGCGTAAACAAATCGGTTTTCGGAGCAAAATACAAAATTCGTAGTTGAGAAAGTCTCTCAATGGCGCA
GTCTGTGATCTTTGGAGTCTCTCTATTCTTTCTGGAACACACTGCTCTGGGGATTCCATGAAGAGACACATTATCCGTGTGCCTTTATCGATACGGCAACATACACGGG
AGCTACGGCTTAGATGGCCCATTTGTGCACAACTGGACGGTGTATCCCGTCACCTTCGTAGCCGTGTACGACTTTGTCTCGAAAATGGCATCCGCATCCCGCAAGCAGCG
ATCTCAGAGCTGTGTCTGTAAGACCAACCCCTGTGTACGATTGTCTTACGCGGAGAAATCTACGACCTGGAGAACAGGAGGAGTCTTAGTCCCTGTGGCGGGGTTGTC
CAGCCTTCCCTCCCATAGCCATATGGAGGTGGAGTTGGGCAACGGCAGTCTGCTCTAGTGAAACTGCAACCCCGTTTATCATCCAGTGGAAACACCATGTGAGCACATG
AAAGCCGTAAACAGGGCTCGGAATACGTTCACTGGACACTTCATGAGTCCGTTTGTGAAATAAACCTTCCATCACATTAATGCTTTTACATCGCCTTTTCAGAACGGAACC
ATCAGCCACCGAGGACATAATTTTCTAAGCAATTACTGCTTCACTCCGCTGCTCCATGGGAACCTCCACCTGGGAGTGGCAGCCATTGGCTGTGCACAGAAAACATATATT
TCGTCTTGGGGTGGGAGTGGACATACGCCATATGTAAGTAGCAAGAGGAAACATTTTCCAGCAACTTCAGCAATTCATTATCCCAATATGCACAATACTGTTTCA
CAACAATACATAAAAGTAATTTGCTAAAGCACCACAAAACAAATCGATGATAGCATTAAATAGTTTGTATAGTCAGAATTTAAATTTACCCAGCGTATTCACTTTCCAGGTCTCTTG
ATAGCTATTGCTGAAGGTTTATCGTGTCTGATGGTCTACCTTATGTTTTCGGAAATCGCAACAGTTTCTACGGTGTGGCCATAAAGCGGTACGCCATTGTATGATCTTGG
GATATGCCCTACTGGCTTATCTCAGCTCCACAACCCAGCAATCTCTCAATGCGGCTTGTCTGATTCTACGTAAGTGGTAAATTTGAGTATTCTGTGATATTTAAACATT
CCCTTCTTTTCTTTTAGCAAGTTTGGCGTTGATGAACCTGGTGTCTTCTTTTACATCTGAGCTTTATTGCAATTAAGCTGTACTTGGCTTTTATGGCGTTGTCTTC
ACAAAATCACTTTTGGCTAAATTTTACTCCCATCGTACTGGTAGCCGTTGGGCTGGTCTTCTTTTGGGTTTTCAGTTATTACGGCTTAGGCTCATCTTCGGAGCGGATA
CATGCTGGTTTGTATGTAAGCACTGCCAATCCATATATGTTTATAATAATTAATTTGATTTCCACATCGCTAGCACGCAATTTGGTCCGTTATGATATATTTTACGCTCCT
GTTTTCGTGGCTTGGCCATTAGCGGTTTCTTCTATGCTCTGAGCCAGATCTACATCCGTGATCAGCCAGATATCGAGACGGGAGAGAGCTTCGAGTCCATAGAGAAGAAC
GATTCAAACTCACTTTGGAAGCACTTTGGCTACACGGCAGTCTGATGGGTCGTTTGCATATGCTCTTTTGGCTTTAACTACTACTGGGAGAACAGATCCCACTTAACTACCG
TGTAAGCTTCTGCATGGCTTCCACGGGTTTGGCGGCTCTATGCGCTGATTGGAAAGAAATCAGCAAAATCCAAATTTTCTGCGGCGCATAGATAAGTAAGTTGATAACTAT
GAATTTTAACTACTCTGTTAAATATATCTGTTGAGTGGCGAAGATACCTGTGAAACTCAGTTCCGCTCTCGAGTTTGGTTAACTTTCTTGTGTGCCCACTGTAGCG
TTAGTGTAACTGGAAGCACTTTGAGCACTTTTCTGAGTTTACGCTTCAACAAATTTATCTTCACTTCTTGAAGTTCCAAATTCCTGGACCGTGATCAAGTTTCTCCAGCTT
GTGTGTAATAATATTTGTAATAATTTGAACCTCGAACAAAAGTACACTTCTACCTTTTAAACGAAATACGATAATAAGATACAAAATAAATATATCTGAAAAAATAA
TCAGGTCTGGAATTTCTACAAATTTGTAATAATGATAACAAGTCCCAACATCTTTTCACTATAGAACAAAACATGGAATATACCTTAAGCCATGTGAAAGAACACATAACATA
TTAATATCTTAAACAATTTGAGCACTTTTCTGAGTTTACGCTTCAACAAATTTATCTTCACTTCTTGAAGTTCCAAATTCCTGGACCGTGATCAAGTTTCTCCAGCTT
ATTTCAAGTCCGATTTTACTGTACCTATCCGCTATCGCCATCGCCAGCCCGGTTTACTTCCCGGATCTCTCTATATTATAAATGTGGCCGATTTTATTAACTCAGAGGTCAA
TGGCCGCAAAACAGACAGAGAGAGAGAGAGAAAGTTGGTCAGAGCCACGAGAACTGCATCCGACAGGGGGCTGTCAAGAGCAAGGAATCAAGTGAATGTTGGTGGTGC
TCTTTTGAAGAACTTTGCAAGCTTTTCCCTCTACACCCCTTTCGCAAGGCTTAATCGCAGAAAACCCCGGGTCCGAGTCCCGCAATCTTATTTACCGGATTGCT
TCGGGTTTGTGCTGCTCTTTTGGCTGGCGCTACAGCAGCAGGACTCGGGGTACACACTAAGAGAGGCTTAAGAATATAGAGATAGACAGGAAAGCAGATGGATCGATA
AAACGTAGTAAATATGCTCGTAGTAATGGAAGCTGGAATAACATGGCAACAAACGCCCCAACTGCATCGACAACACCATCAATCCAGACGTGGCCACTGGAAATCC
CGAGAGGTAGGACATGTTTGGCGGTTTTCGACTGCATGTGTCGAGTGTCTGTTTGGTTAGATACGTATGAAAAGTGAAGAAAGCTTATGACCGCGCCGCTAGTA
TCGCATAGGCAATATCGATATGGGAAATCATTGACTAACATTTGCAGACATTTAGTTTGGGACACATTAAAGTTAAATTTGACTC
(SEQ ID NO: 145)

Exon: 1001..1504
Exon: 1560..1716
Exon: 1897..2088
Exon: 2149..2367
Exon: 2427..2784
Exon: 2838..3116
Start ATG: 1001

Transcript No. : CT32762

ATGGCGCAGTTCTGCATTTCTGGAGTCTCTCTATTCTTTCTGGAACACACTGCTCGTGGGGATTCCATGAAGAGACACATTATCCGTGTGCCCTTTATCGATACGGCAACAA
TAACGGGGAGCTACGGCTTAGATGGCCCATTTGTGCACAACCTGGACGGTGATTCCCGCTCACTTCGTAGCCGTGTACGACTTTGTCTCGAAATGGGATCCGCATCCCGCC
AAGCAGCATCTCAGACCTGTGTCTGTAAGACCAACCCCTGTGTACGGATTGCTGTATTACGGGGAGAAATCTACGACCTGGAGAAGAGGAGGAGTGTAGTCCCTGTGGCC
GGGGTGTCCAGCCTTCCCTCCATAGCCATATGGAGTGGAGTTGGGCAACGGGAGTCTGCGTGTAGTGAACCTGCAACCCCGTTTATGATCCACGTGGAAACACCATGTG
AGCACATGAAGCCGTAACCAAGGGCTCGGAATACGTTCACTGGACACTTCATGAGAAGGAACTTCAATCGCAGAAAACCTATATTTCTGCTTGGGGTGGGGAGTGGACATACGCCATATGCTCTTGATA
GCTGCTCCAATGGGAATCCACCTGGGAGTGGCAGCCATTGGCCTGTGCAAGGCTTAATCGCAGAAAACCCCGGGTCCGAGTCCCGCAATCTTATTTACCGGATTGCT
ATGCCCTACTGGCTTATCTCAGCTCCACAACCCAGCAATCTCTCAATGCGGCTTCTTCAAAAACCTGATGTTTGGCTAAATATTACTCCCATCGTACTGGTAGCCGTGGGCTGGCTTTT
GAGCTTTATTTGCAATTTAAGCTTACTTTGAGCTTTTATGGGCTTGTCTTCAAAAACCTGATGTTTGGCTAAATATTACTCCCATCGTACTGGTAGCCGTGGGCTGGCTTTT
TTTGTGGGTTTCACTTATTACGGCTCTAGGCTCATCTTCGAGGCGATACATGCTGGTTTGTATCCAGCAATTTGGTCCGTTATGATATATTTTACGCTCTGTTTTCGTGG
CTTGGCCATTAGCGGTTTCTTCTATGCTGAGCCAGATCTACATCCGTGATCAGCCAGATATCGAGACGGAGAAGAGCTTCGAGTCCATAGAGAAGAACCGATTCAAAT
ATTTTGAAGTACTTTGGCTACACGGCAGTCTGATGGGTCGTTTGCATATGCTCCTTTGCTTTAACTACTACTGGGAGAACAGATCCCACTTAACTACGCTGTAAGCTTC
TGCATGGCTTCCACGGGTTTGGCGGCTCTATGCGCTGATTGGAAGAATCAGCAAAATCCAAATTTTCTGCGGCGCATAGATAAGTGGCAAGATACCTGTGAAAACCTCAG
TTCCGCTCTCGAGTTTGGTTAACTTTTCTGTTGTGCCCACTGTAGCGTTAGTGTAAATATATTTGTAATAATTTGAACCTGGAACAAAAGTACACTTCTACCTTTAAAC
ATCAGCATAGACAAAACCATGATTTTAAATGTAATAATTTGAGGATGTGTGTAATAATATTTGTAATAATTTGAACCTGGAACAAAAGTACACTTCTACCTTTAAAC
GAAATACGATAATAAGATACAAAAT
(SEQ ID NO: 146)

Start ATG: 1

MAQFCILGVLLIYSGTHCSWGFHEETHYPCAFIDTANITGSYGLDGFVHNWTVIPRHFVAVYDFVIENGRIPASRLHACVCKTKPCVRICCLRGEIYDLERQCLVPVA
GVSSLPSSHSMVEVLNGSLRLVKLQPRFSIHVETPECHMKAVTKGSEYVHWLHENGTSIHRGHI FSKHYCFTPLHGNSTWENQPLACAPKLYFVLGVREWTYACILLI
AILSMFIVLMVLMCMEMRNSFYGVAIKAYAI CMLGYALLAYLTLHNPANLSNAACRILPSLALMLNLSFYILSFTAFKLYLSFYGVVFTKLFWLFTFIVLVAVGWSF
FVGFSGYSGRLIFGGDTCFDPDRNWSVMYIFYAPFVACISGFFYVLSQIYIRDQPIETEKSFESIEKNRFSFKWYFGYTA VVVVVCICSFAFNYYWENRSHLNVAVSF
CMAFHGFAALYALIGKNQIQNFLRRINDGEDTCENSVP LSSSF*
(SEQ ID NO: 147)

FIGURE SHEET 65

Name: mth-like 8
Classification: G_protein_linked_receptor

AGTCAGTGCCTGGACGGATCGCACGAGTCTTCTACTAAGCGCTACCAGTGGTGGTCGTAATCCCCCAATCCAACCCATACCCAAAGCTTCGCTGCATTTTCGAATACAAAC
CACTCAAGTAATTCCTCGCTGGTGAAGACTACCTTATCCATCAGGATTACTACATATCCCTACAAAAAATATGCTGGTTAAAGACTATACACCTAGCTCCATATACATATATAT
ATATAACAATATCGAGGCATGTGTTGTGGGCGAGTGAGTGTGCGGTGACTCTTGATGTAGTTGCAATTTGATTCGAGTATTTTCGAGTGCTTAGTTTGTAAATTT
GTAGCTAGCCGTTAATGTTTGTATAAATGTTTCATTGGTTTCCCACTCCGCTTTATTCGATAAGCTCGTTTTCTAGATTTTAGCTTTAGCACTCTTAGTTTAATGTTTAA
GTTAGTTTCGCTGTTAAGTTTACACCGTAGTCGAGGCGCTCTTGAGTAGTCTCTTTGTACATAAAGCTTACACTAGCTCTACATAGATATACCATGTCGAACCTACGAGTC
TACGAAATACGAAACATATATGTGCGTTGAAAGCGTTGTATTTTGAAGCGGAACTTATGCAAAATGCAAGGACAATAAACAATATGCTATAAAGACGACCAAGTGTCTC
GCAATAAATGTGAGCCACGCTTAAGAGAGACGATTGTGCGCAAGGTGTTTACTCGCGCGCGCGCGGCAAGCGTTTGGGGCAAAACGACCGTAGTGATGTTTGCACT
GTACCATCATCATGACGATGGCGTGTGAACGACGCTGCGCATAGTAATGACAGCGGATGGCAATCTGAACCATAGCGCGCACCCGATATAGAAACATTAACCTAAACCGTCG
GTAGTCGCTGGCCATAAATCTGTCGGCAGCTGCTGCGAGAAGCTTTAGGTGCGGACGCCCTTTCGCGGCATACTTTCGGGCTTCGGTTCGGTTCGCCAATGGCCCT
GCTCCACTACACCTTCGACAGCGTCAAGTTGTACTAGAAATGGGCGTTCGCCAGCGAGAGGCGACCGCGATTCCGCTCGATACGCCGTAATCCGGGCGTGTGTCGGGCG
GATCTCAGGCAAGCTGAACCGCTTCAAGCGACATGCCTTCTCCGCGCTGCTGGGCAACGCTCTTCGTCCTCGCCTCTCGCGCAACCTGAGCACCCCTCTATGTGAACAGCGCGC
GGAAGCTGCGGCGCCTTCTTCGCGGCCCTGCCATCTCATCTGCGCTGCAAGTACCTTGTCTCCGACAATTTCTGCACCGTATCTCTACATGGCCACAGTTCGCGCGCATCTCT
GCACTCTGGGTGAGCGCTGAGGGAATTGAGCAGACTCAACAACAGCTAATTAACAATCTCTTAGACAAATTTGGTGGCTTCATGTGCAAGTTTGTGGCGTTCATAACACCAAC
GTCGGTGTGTCGCGGACGCTGACCTCGTGGTGGCATGCGCTTGGACCGCTACTTGGCGCTGATGCGACCCGTGCTGGGGTCTCGAGTCCGACAAACGCTTCAGCACTCTG
AGCATGCTGCTCATCTCGGGATCTTCGATAGGATCTCTGCGCCGCTGCTGGGCACTGAGCACTACCGAAAGATCTATCTCTGGATGTGGAGAGCTCCAGCGAGGAGACGG
AGAGGTCGTGACCGCTGTGCCGAGGAGCTGGTGGTCACCGACCTGGAATGTTTACATGTGCTGGCGGGGACGTAGTCACCAAGCATGGATTATCGCTTAATACAGTT
GACAGCGATTAAAGGGATTTGGACGGGGGATGTCAAGTTGTTGAATCAATGTTCCGATATCTTATACAGCCGCTCGAAAAACAATTTATGCAATCTATAAATATCCGTT
TTAGCAGCAGCTCGGACTCTACTACGTCATCTGTTACCTCTGCGGTGATGTGTCGTCGATGTTGCTTCTCTATGGCTGAATGCGGTATGCGGATGTGGAGGACGCTTCGGTGGCG
AGACATACCAACGAGGACGAGGACGAGACGATCAGGAGCCCAAGGGGTGAGTTTAAACCATGGCCATGCGGGGATCTTGTATGCCCTCCACCTTGGTCATGTGCA
TGGGAGTGGCGCTTCCCTTTGCGCTGGACAACTCTCTTGGCGCCAAGAGTACCGTCAACGCTCTTCGGCAAGAAGACCACTGCTGCTGCTCTAGCCAGGGAGGCCAGGCA
CCCGAAGTGGTGTGTTGGTGGTCTGATGATGAGGCGGTTTTCATATGCCCTTCGACTGCGCGCCTGGGTCTTCCTGATCATGCGGCTTTATGGAGCTCTACTCCGAGCCATC
GACGTGGCTCTGTGCTTACGCTTCGCGATCTGCACTAGAACCTTTTTCATGTGCGCTCAATCCCATCTTCTACACCTTCTCATCCGACCATTAGGACCTTGACGCTGGTGCAT
ACAAAAATTCAGGGATTTCTTGGCTGTCGCGCTGTGAAGTGCCGGATGGTATGCCACGGATCAGATGGACAAAAGCGGGTGCTGCTGCGGCTGAGGCCACCCACCTTCAC
ATGGAGGTGCCATCCGAGCGGATAGGCGAGCAGCACCGTATTCGGGATGTGGACAGCCGATCCGCGCTCCGACAGGCTGCAACCCGATCCGAGTAGCTTACGCGG
TCTCTCAGCTACAGCAGGAGGTTTACCATTTACAAGACGTCGGTATGCTCTGAGTGGCTCAATGAGTCTCCGCTGATAGCGGTGGTCTATAAATCACC CGGGT
CATTTGTGCGGATACGGGGGCCAACGGTTAGCAGAAACCCAGAACCTTGATTGATGGATGGATCCAGCCCCCATCTCATCTGTACAGGCGCAAAAACGGTTTGTACGGTGGGT
ATCATACCCACGAATACATACGACGCAATATAAGATATAAGCTACATATATGTACCTCATCAAGCTGTGAATATAGAAGAAATACGAATGGGTGTTGGCAGCTAACTAA
CCATTAAACATATGGCGCAATCGAAATCGGGAAGGTCTCAGCTATTATTGTGACGAAACCTTTAAAGTACCACTTAAACATATATTAGAGAACTTAAGCAAAAT
GTGAGATTAAATAGCAATGCAACTCTTAAAGGTGCAATATACCTATTTCTTAAATATTTTACTTAGAAATATGACATCTTATATGTGCTTAGTTTTCGTTACAGACAGT
ATCTTTGTATTTCTAAGACGTAGTAATTAGTTAATGGAAATTAATTAGTAACGGAACCTTAAATTAGTTCATGCGCAAAAAAGAGAAATGGGCTGACAGCAGACATTA
TAGAGTCAGCAGACTGCTGGACGAGACAATGAGCCCGCAATGCAATGGAATAGATTGATCATATATAGATATCTCCGCTTGGAAACCAATGGGTTAAGACAGATCTCGTGAGAGA
TGGGGTGAAGAGCTGTTAATTTGTCAAATGTGAGACGCGGATTAATTTAGCTTAGCGAGGAAAGCTCTCCCATACTCTCTTCCATGAACTTTTAAAGCCGACG
AGCCCCAGCTCGACTCAAAGCTGAAATTCGCGTTCCTGCAATTCAGTGTGCGCGGTTTGGCGGACGCGGTTGGATTGCGATTCTTGGTGGTAAACGCGAAATCTTCT
GTGCGCGGGGTGAAATCTTCGCGAGAGAAACAACAAAGCAGCAAGTGTGGAGTGGATGTCATACCGTACACG

(SEQ ID NO: 148)

Exon: 1001..1354
Exon: 1410..1757
Exon: 1909..2772
Start ATG: 1001

ATGGCCCTGCTCCACTACACCTTTCGACCAGCTCGAGTTGTACCTAGAATGGCGCTTGCCCGACGACGGAGAGGCCACGCGATTCCCTCGATCCAGCCGTATCCGGGCGGTGT
TCGTGGGCGCTCTCAGCCAGCTGAACCCGTTCAAGCAGCAATGCTTTCCGCGCGCTCGTGGGACGCTCTCGCTCTCGCTCTTCGCGGCAACCTGACACCCCTATGTGAA
CAGCGCCGGAAAGCTCGGGCCCTTCTCCGGGCTGCTCATCTCACTGGCTCGAGTGCAGCTTGCTCCAGCATATTCTTCAGCAGTATCTTACATGGCCAGTTCCAGCG
CAGTATCTGCAGCTCTGGACAAATTGGTGGCTTACATGTGCAAGTTTGTGCCGTTTCATAACCACCACGTCGGTGTGTGTCGGCAGCCTGACCTTGGTGCCATCGCCTTGAGCC
GCTACTTGGGCTGATGCGACCCGTGCTGGGGTCTGGAGTCCCGACAAACGCTTCAGCATCTGAGCATGCTGCTCATCTGGGCAATGTTTCATAGGATCGCTTGGCCGCGCT
GCTGGGCATCTAGCATACCGAAAGATCTATCTCTCGGATGTGGAGGACTCCAGCAGGAGAGCGAGGAGGTCTGACCGCTGTGCCCGAGGAGCTGGTGCTTCCACCGAGCT
GAAATGGTTACATGTGCTTGGCCGGGACACGACGTCGGACTCTACTACGTCATACTGTTACCCGTGATCTTCTGCGGTGCATTGTGTCTTTCATATGGCTGAATGCCG
TGA TTGCGAGGACGACTCTGGCTCGGGACAGCATACCCAGGAGCAGGAGCAGCATCAGGAGCCCAAGGAGGCTGATTTAAGACCATGGGCAATGGCGGGGATCTCTT
GATGCCCTCACTTGGTTCAGTGCCATGGAGTGGCGGCTTCCCTTTGGCTGGACAACA CTCCCTTGGCCGCAAGAGTACCGTACAACGCTCTGGCAGAAAGCACTGCT
GCTGCTTAGCCAGGAGGCCAGGCACCGCAAGATGGTGGTGGTGGTCTGCTAATGATGGCGTTTTCATATGCCCTTGACTGCCCGCTGGGTCTTCTGATCATGCGGC
TTTATAGGATCTCATCCGAGGACATCGACTGGCTCTGATCTCAGCTTCGGCATACTGAACCTTTTCACTTGGCGCTTAAATCCCATCTTCTACACCTTCTCATCCAGAC
CATTAGGACTTTGACGCTGGTGAACAACAAATCAGGAGATTCTTGGCTGTCCGCTGTGAAGTGGCCGATGGTATCCACCGGATCAGATGAGCAAAAGCGGGTGTCTG
TGGCGCTAGGCCACCCACCTTCATATGGAGTGCCATCCAGCCCGGATAGGGCAGCAGCCACCGTAATCCGGGATGTGGACAGCCAGATCCGCGCTCCGACCAGGTGC
AACC GGATCCGAGTAGCTACGGCGGTTCTCATAGCTACAAGCAGAGGAGTGTTCACCATTTACAAGCATGGCGGTGACTCGTCAGTGCCTCAATGAGTCTCCGCGCTGA
(SEQ ID NO: 149)

Start ATG: 1

MALLHYTFDQLELYLEWAFAQHGAEATPISIQPYPGVFVGDSLQNLNRFKRHAFSAVVGTLFVLAFCGNLSTLYVNSRRKLRPFFRACLISLACSDLVSSIFCTVSYMAQFQA
QYLQWLITGGGCMKEFVPIITTSVLGSLTLVAIALDRYLAVMRPLVGFWSGPKDRFSTLMKLLIWAIGSSGPGLLGYDYKRILVLDVDESESEEVVTAPEELVTEL
EMVHCLAGDHDVGLYVYLFTLSFLCPIVSFLVWVARLOGLWRRHYHQEQQHQEQKEGQFKTMANGDDLMPSTLVSAMGVAVPFALDNTPLPKXSFTVNAPEGKTTA
AALAREARHRKMVVVVLLMMAVFICLRLPWVFLIMRLYGSYSEPIDWLLYFSFGILNLFSCALNPIFYFTFLTQTIRTLTLVKHKIQGFLGCPPGKVPDGMPTQDMQDSGCC
CGLRPPTFTWRCHPSRDRAAATVIRDVDQDPDPSDQVQDPDSSLRRFLSYKQEVFTIYKQCGDSSSASIESSA*
(SEQ ID NO: 150)

Name: Neuropeptide receptor-like
Classification: G protein linked receptor

FIGURE SHEET 67

GCTCCGCTCCTCAACGCTGTGCATCTCGTCTGGAGCTGGCCGCTCTTCTGGGCTGCGGATGCGCTCAGCTACTCCAATCGGCCATGAACCAATACATGTACGCCCTTTTGAGCGATAACTCTAAGAAGAGGCTTCAAGGCGCTCAGCGTGTCTGCCGCGCAAGGATGTGAATGCCCACTTGCAGCTGGAGAACAGTTCTTCCCCAAGTTTGGCAGAGGGCAGCAATCGGAGGCTCTTCTTGTGGCAATGGAAAGAGTGGCGGCTGCGCTGGGGCATTAACCAAGAAGAGTACTGCTGGCGAGCAAAACAATCTCCGATGGCCACTACAAACGACGACGACCAACCAACCGGGCAGGATGTCAGTACCTGTCTCAGCCGCGGATACACAGGATGCCACGGGAGATCCAGCTGGGAATCCGGCCGCGGCTGCTGGTCAATGTGAGCCAAACAACCTGTAACCCGCGGTGCTCCACAGGCACTTAAGACCGGGCTCCCTCGATGCCCTGGAAACGGTGGTCTTCTATCGCGGAGGATGAGACCATCAGGTTGAGCTTGGACCTCGATACGGCCATCGATTGCCAGGCAATAGCGAGGCAGCCGGAATGCTTGCTATAGTTCGCCATAAATGATATTACATTTTATACCAATATGCGAGAGCTTTGCCAAAACGGGTAAACTTAATCCCATTTTATATGATCGAATCTCAGCAAGAGAAATTTTAAAAACCCGGAATAAA
(SEQ ID NO: 154)

Transcript No. : CT33159

ATGGAAGCTGGATGGTGGCGAGGAGGAGGAGGAGGAGGAGCGTTTGGCGGGAAAGCCATCATGGAAGGACACTCGACACCAAAATGGGGCAGCTGCAAGCCACAGAAACAAC
 GCACAAGGACCAATATCGCGACCAACGGCTGTGCCCATTCGGGCATCTCGTTATTGTGCTGACAGCTATGACATAAGCGCTTAATAACAGCCGACAGGAGCTGGCAGT
 GGGCCCAAATGGCACCACTGCATCAACTGGATCCGTGGAGTCCGAGTCGATATCCATCATTAATGGCATGCAGATCAGAATGAACAACTGGTGACCAGTGTGGCGACCCCACTT
 GACCAACGGAATGCACCAGCAGCAGCAAGATGGCAGCTACATATTGGAGTACGACAGTATGGCCCCGAGCTGTTCTGTACAGCTACAACCTTCATCCTGAAGCTCATCACGATGA
 TCCTGTAGCCTAGTCTGTCATCATTGGACTCTTTGGCAACACCCGTGTGATCTATGTGGTATGAGGTTCTCCAAGATGCACAGCGTAAACACCATATACATCTGAATCT
 GGCTATCGCGGATGGTGCTTCTGTGATCGGCATTCCTCTGCTCTACACATGCAGGTGGGCAACTGGCCCTCTCGCAACTATATGTGCAAGGCTCATCTGTGAGCACT
 TCGATCACTCTCTTCACTCTCTCGATCTCTCTGTGATGTCGCGCGAGTCGCTACATAGCCGTTTGGCAATCCCATATCTTCGCGCTCGCTACCGAACGCCCTTTGTATCCA
 AGTTGGTTTCGCGCTTCGCCGTGATGACATCCGTGCTGCTGATGCTGCGGTTATCCTTTTGGCAGCACCGGTGCACTGCAAGCAACGGGAATGTGCTCTGCAACATCGAGTGT
 CGGCACAGCTCAGAATCGCACACCGACTCCACCTTCTATTTTGTCATCGCTGGCTTGGGATTCGCCACTCCACTGACTTTTATCTCTGTGTTCTACTGCTGGTGATCAGG
 AAACCTCACACCGTGGGACCGAAGCAAGCTTAAGGAAGAAGCGCTCTCACAGGAAGGTCACCAAGTTGGTGCTCACGGGTAATAATATATGACATTTTCGATTAGTTGTG
 GAACTAACTACTACCTGACCACTCTGTGTTCAAGTTTGATAAATATTATTATACATTATAAATATATATTTTTTTTGACATTTTGAACTCGACTATAAATTAATGTGTTT
 AAATTTGATATATAGAAAGAAATTTTATAATATTATTCAATACAATTTATGTTTTTTTGATTTCTCCACAGGTCATAAGTGCGTCACATATTTGTTGGCTT
 (SEQ ID NO: 155)

Start ATG: 1

MEGGWRRGGGGGRLGGKAI MEHGSTPNGAAASHRRNNSTRNIATNGCAHSGILLFVL TAML TSLTIPTEQLAVAPNGTTLHQLESVESESYP SINGTQNETMVTSVRPHL
DHRNRPTQONGSHYLEYDDG PDCSYSYNFILKLITMILYALVCII GLFGNTLVYVVMRF SKMQVTNIIYLNLAAI ADECF LIGIPFLLYTQVGNWPGNGYMKAYMVST
SITSFTSSIFLLIMSADRYIAVCHPISSPRYRTPFVSKLVSAFAMT SVLLMLPVLFASTVQSSNGNVSCNIEWPDTQNSHTDSTFILYSLVLGFATPLTFILVFYCLVIR
KLHTVGPKHKHSEKKRSHRKVTKLVLTVIIIDISISCGTNYLTLTVFSLINIITL*
(SEQ ID NO: 156)

Name: Somatostatin receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013383997

GenBank accession number: [F42800.1](#)

CAATAAGACCATTCACATTTCGAGCTGCGACGATCGTATCATCAGTTGTTTGCAGGACCTTGAGTCTCTCATACAAGTTTATAGCTGCTCATTTTAAATCTGTTCTTGTCGCA
GAGATTACACCTTCTCCGAAAGCGGCGACGACGTAAGATTGGAAAGAGGGTTAGTGCAACTCTTTGTGCAATTACAAATTCATACATACCGCGTAGTCTCTCTCTGCGTCGACAA
GAAAGAAATTGGTATGTGGCGCATGCTGTCGAATGCCATGCAATTCATGCCCAATCAAATAATTTGAAATGTGTAACAACACTCACTTGTTTTCTTTCTGCGGCGCATGTGGT
TGCTCTCGGCCCATTTGCCATTTCGCCGAGGCATAATCCCGACAATCGGTGCGATGTGGGTGATACGAGTGCGAATGCGAATGCGAATGCCAATGATATGTGTATTGTATGTAT
GTGCATACATATTGCGCTCGTATGCTGTGGGCTGATGTTTGCACTCCGGTGCAGCGGAGTATAATAAGCCGGGTATTTATTATGGGCAATCAATAACTCGGTTAACGACG
GGCACTCTGCCAACGAGATAAGATCTGCTGTAACAAAGGCAACCGGTGCAGCTACGTTCTATTGTGCGAATAGCTTTCAATTAATCTGACCCGGATACCGGCGGCGCATGGGT
CGGGGAAATCGGAAATCGAAACGAAACGAAACGAAAGCGGAAATCGAAATCGAAATGGAAGAGATCAATTTTCAATTGCAGCCAATGCATGCATGTCCATGGGTCCCA
TGCTAATATCGAAGACAAAGCTCTGATCTCTGAGGTCGTGTGCCATCGCAGACTCATTTGGTTGGTGGGTTTAACTCGTACAAGATTGTATGGGTTTGGGATGTTT
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CCTGTCGAACATTCTGACTGCGGAGGCTGTATTTCGCCCCAGCGCTTCAACCCGCTGCTGCGCAGTCCGCGCTCATCAGGCAATCCCATCTGCACGACGGCGGGAACA
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TCCTCGACGGTGTGCGAGGGTGGCCCAATTGGCCCTGGAGCCACCGCTATGAGACGCACTGCGCGATTGCGACAGGAAAGGTTTCGGTAAATCAACAGCAGCGGCC
CTGGAATCTAACGCTTGCGTCGGCAGCGGCCACCAATTCGAGAACTGCTCGCGCTTTTGTCAATTACACCTTACACAACAGGTAACTAAAGCTTAAATAACTTTTAA
TCATATTGCGATATTTTTTTTTTTTTTTTTGTGTCGGCGGATAGGACTACTGTAATGGACCTGGGACCAATGCTCTGATGCTGGCCACCCACTCGGGCTGGAGTCTTGAC
GGATGAATTTCTCTGGCGGCTTTTATGCGGTAGTGGCGGACGAGTAAGTAATGAAATACACTCTAAAAAAAAGAGCGCCGAAAGAACCAATGCTGAGTACAATTAGCATAT
TAATCGTTGAAGCTGTTTAGCTGTTTAGCAGCCACAAGAAGCTTATGACTTCTGCCGGCTTTATTAGGAGTACAAGACATGTCTTACGGGTGTTTATCCATCCACACGGCA
TATCTGGCGGCGCAACATACCGGATATTAGCTGGGAAAGCGTTCAAGTGGTGACGGCAAAACAAATTTGATTCGTTATCATCGAAAGGATGCTAATATTACATGCA
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ACAGGGTTTAAACACCGCAATCGATTGCTTAATTACGATCTTAATGCTTGCTTACCCTCCGATAAATTAACAGCTTCTGCAATGGGGAATGCTATTATTAATCAACAT
GGGCGCATACATACGAGGCGAAGAGTTCTAAGTAGCTACTTACCACCAATTTTACAACTTAAATGCTCTTCCATTATCACTAGTGAATGATTATCTCGGCTGTGTG
GTGCAAACTTAGGCATATGCAATATATCTGTGTAGAAATAAATTAATGCTAGTGCATTTGTGAGCTAGCGGCTATTCTCGCGGCGTGTAGATTTTAGGCCCAATAT
TTTTCAACTGGACGAGGTGGGICGCCCATTCGATTGGCATAATTGCACTTGTGACGTCAAGTGTGCAAGGTGCAATTACGTCATATGTAACAACCAACCTCCCTCTTTTCCGG
TTCGAGAATTCGCCCATCCGAAAGTGAGCGTGGAATGCTGATGGGGCAGCGGCCAGTGCACGAGGTGAATCCGCGGATAGGACGGACTACGGGCGGTGTACAGCT
CGGAGATATTCCGCTCTATGCAAGCATGTGGCGCAAGGACTTCGATGTGATGAAATCTTTTCCACATCAATCATATAACTGTTTCAGTTATAGTAAAGCATTTGCTT
AAATGCCAACCAAAATCCCAATCAAGGGTATGGTTTTGAAATTTGCTAATAGGGTTTCAAGTTTTTTAGACACCCCTAAAAAGTGATTTCAAAGGCTGATTTACATTTTGTG
ATAGTGGTTTTATAGTAGTATTTTAAATCTCATTTTCTTCAACTGGTTTTGCGATATCTTCCGCTACATAGACATTCGCGGAGGACTCGAACCCTGGAGATCTGGGCT
CTGCTCTCTCCCTGTTCGCCCTTATAGTTTCCCTGCTGATCTTCTGCACATTTCTGCTCGCTGGCAAAATCGCACCAAGATCACAGAAATCTTTCTGCGCATGGTGCTG
CAGGTGATCATTGCGCTGACCTTGATCTCGACCAATTCGGCGGGGAAACAGGAGCGGCCCAACACGAGTCTCTCTGTCTATGAGAAACCGGTAGCTTAATATTTCC
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ATGTGGAATTTTACATCGAGGCGCTTACATGCAACATGCTGACCTGGCGGTTTTCCAGGCGACTTTCCCTCAAGTTCTTCGCGACTCGGCTGGTGTGCGCCATTC
TGATGACCACCGTGTGGCGAGATGCAGGTCATGATATGACACATCTCGTGGGCGAATGCTTGGAACTATAATCTCAGCCCTACTACTGGAATCTCTGAGGCGCATTC
ATAGCGGCTACTACTGGTGGGTAACTCAACTGTGATCGCTTAACTCACTATTCACCACTGAAACAGTCCGATGCAGCTAAACTCTGTTTCTGTTGAACATTCAC
CGATGTTCTGTAATAAGAGCTGCGTCAATCGCAGGCGACGATATAGAACAGACTGCAAGGCAAGTAGAGCGGCTATAGTCTTACTACACTTTTGGGTATGAACCAATCTCC
TGCACCAGCTGGCTCTCTGAAACCGGCCACGAACTTCGCGGTCTGTGTCGATGGCACCACCTTCTCACTCTGTTTCAGGGAATTTTATAGCGCTAATTACTGCTTTCT
AAATGGCGAGGTAAAGCATCAAGTCACTCACTGATTTGCTTTCCTTAAACCCCTCACTATTCCAACCAAAATATATTTTTAGTTTGTGCGCGTCTCACTAAAGAGTCTC
GCCCTAGGCTGTGGTGGCAGGTCATCGAAATGGGCGCGAAAGGCAATCATGTAATCGGGTGTATAACACGCGCGCGGATACGGATGCACTGAGCTGACGAGTGCAGGAG

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ATCCATCGGCCACTGGAAGCGGTAGCATCCGCTTTGTGCACACCCACCTTGAACTCCTTACATCAATTTAAACTCCCAGAATATCACCGCCGAATAAAAGGCTGAAT
GGAAGAAAGCCGAGCAGTGCAGCATTTGTAGTATTCAGAGCCTCAACAGCGCCAGCAGTGCATGCCCCGGCTGCAAAACAGGCGCGGGAAGGGCAAGGACCGGGTGG
AGAAAGCGGATGCGGAAGCGGAGCCGATCCGACCATCTCCACATTCACAGCAAGGAGGCGGCGAGCGGAGATCGCGAACTCGCGCTCCAAGTGGAATAAGGGCATCTG
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GAACAGAAAGTAAGCTCATTGTTTTGTAATCAAATTAACCAATAGACATTAATTTAAATAGTTATTTTCGCTTGCCTTCGGTCTTCTGCTCCACTGTACTTATGTAATA
CATACATACCTTTATTTGTGAAGCCATATCCAAACCTTTCCAAATCTAGAAATCTCCTATCCTTTGAAATCGGCTAATCTACGAAATTAATTTCCCTACAGTT
TTACAAATATTTTGTAGACAATAAATTTCCAATAGCGTCACCTTCTGTTCTGGACACGCAATACATTCAAAACACTCGCACACGGTCTTACTCGATGAATTTTTCAG
CAAACTTCAGCGGCATGTTCTTTACTTCTACTATTATTAATATTAGATTGAAATATTTAAGTGAATTAACACAACTTGGCGTAAGTGCAGTGTGGCAGGCT
TTGTAGCCATTAGAACCTATTACATAACATAATTGAAAGTGAATTGTTTAAAGGAACTCACCTTTAATCACTGTTGAAGTGTGCTGTTATCGGATTCCGTTATTAAGT
TTTAAGATTGGGTTTACTGTATTACATTTCTTTACCATTGATTAGTTACACTTCCGTTCTGAGCTGGTCACTGCCAGCTACAATCGATCAGTTCACTGCA
ACTGCCTTGAATTTTGCTAGCTGCGCTTTGTGTGCAACAAACTTTGCCACAATCGATATTTAACTAATGGGGT
(SEQ ID NO: 157)

Exon: 1001..1322
Exon: 1500..1610
Exon: 2472..2623
Exon: 2863..3120
Exon: 3192..3488
Exon: 3555..3818
Exon: 3894..4054
Exon: 4117..4453
Start ATG: 1001

Transcript No. : CT33238

ATGACCTCCTGTCGACATCTCGACTCGGAGGCTGTATTTCCGCCAGCGCTTACCCCGCTGCTGCGCAGTCCGGCTCATCAGGACATCCCCATCTGCACCGACGG
CCGGAACATTTGAATCAAAATCCATGCTGGAGCCAACATCTCGCACAGCCTGGCGACCGGACGCTGCCACTACTGCACGATTTTCGATGCTCGACACCGAATCGCCGGG
AACCTATGCTCTCGACGGTGTCCGCGAGGTTGGCCCAATTTGGCCCTGGAGCCACCGCTCATGGACGACGCTGCCGATTCGGACACGGAACAGGTTCTCGGACTCTACTGCAAT
TGGACCTGGGACACATTTGCTGCTGCGCCACCCACTCCGGCTGGAGTCTTGCACGGATGAATTTGCTGCGGCTTTCATGGCGTAGATACCGCAAAATTCGCCATCCGAA
AGTGTGAGCTGGATGGTTCGATGGGGCAGCAGGCCAAATGCCAGGAGTGAATCCGCCGGATGGACGAGTACGGGCGGTGTACAGCCGGAGATTATCCGTCTCATGCA
GCAGATGGGCGAGCAAGGACTTCGATGCTTACATAGACATTTGCCAGGAGGACTCGAACCTGGAGATCGTGGGCTCTGCTCTCTCCCTGTTCCGCCCTTATAGTTTCCCTGCTG
ATCTTCTGCACATTTTCGCTGCTGCGCAACAAATCGCACCAAGATCCACAAGAAATCTTTTCGCGCATGGTGTGTCAGGTGATCATTGCGCTGACCTTGTATCTCGACCAAT
TCCGGCGGGGAAACAGGAGGCGGCCACCAACACGAGTCTCTCTGTCATTGAGAACACGCCCTATTTGTGCGAAGCATCTATGTACTTCTGGAGTACGCTCGTACCGCAT
GTCATGTGGATGTTTATCGAGGCGCTTTACCTGCACAAATGTTGACCGTGGCCGTTTTCAGGGCAGCTTTCCTCCTCAAGTTCTTCTCGGACTCGGCTGGTGTGTGCC
ATTCTGATGACCCCGTGTGGGCGAGATGCACGGTCTGTATATGGACACCTCGCTGGGCGAATGCTTGTGGAATATAATCTCACGCCCTACTACTGGATCTCTCGAGGGG
CAGGACTAGCGGTCTACTGCTAACTTCTGTTTCTGCTGTAACATTATCCGAGTGTGTAATGAAGTGTGCTCAATCGCAGGCCAGCGATATAGAACAGACTCGCAAGGC
AGTTAGAGCGGCTATAGTCTTACTACCACTTTTGGGTATAACCAATCTCCTGCACGAGCTGGCTCTCTGAAACCGGCCACGAACTTCGCGGTCTGGTCTGTATGGCAGCCAC
TTTCTCACCTCGTTTTCAGGATTTTATAGCGCTAATTTACTGCTTTCTAAATGGCGAGGTTCTGCGCGTGTACTAAAGAGTCTGGCCACCCAGCTGTCCGTCGAGGTC
ATCCGGAATGGGCGCGGAAAGGGCATCTATGTACTCGGGTCTTATAACACGGCGCGGATACGGATGCAGTGCAGCTGCAGGAGATCCATCGGCCACTGGAAGCGAAT
ATCACGCCGAATAAAAGGCTGAATGGAAGAAAGCCGAGCAGTGCAGCATTTGTGATGATTCACGAGCTCAACAGCGCCAGGAGTGTATGCCCCGCTGCAAAACAGGCG
CGGGAAGGGCAAGGACCGGTCGGAAGACGGATGCGGAAGCCGAGTCCGACCATTCGCCATTACAGCAAGGAGGCGGCGAGATCGCGAATCGCG
GCTCCAAGTGATAATGGGCATCTGCTTCCGGGTCAAAGGTAAGAGTACCGTCAAGCTCATCCGTGCCACCCGAGTCACTGTATTTGAGTTGTGAGCAGTAG
(SEQ ID NO: 158)

Start ATG: 1

MTLLSNILDCGGCISAQRFTRLRLRQSGSGSPSAPTACTFESKSMLEPTSSHSLATGRVPLLDHFDASTTESPGTYVLDGVARVAQLALEPTVMDALPDSDEQVLGLYCN
WTWDTLLCWPPTPAGVLARMNCPGGFHGVDTRKFAIRKCELDGRWGRPNATEVNPWWGTDYGPYKPEIIRLMQMGSKDFDAYIDIARRTRTLEIVGLCLSLFALIVSL
IFCTFRSLRNRRNTHKLNLFVAMVLQVIRLTLYLDQFRNGKEAATNLSVIENTPYLCEASYVLLLEYARTAMFMWFIIEGLYLNHMTVAVFQSGFPLKFFSRIGWCV
ILMTTVNARCTVMYMDTSLGECLEWNLTPYYVILLEGRLAVILNFCFLVNIIRVLVNMKLRSQASDIETRKAVRAAIVLLPLLGITNLLHQLAPLKTATNFVAVWSYTH
FLTSTQGFIALIYCFNLGEVRAVLLKSLATQLSVRGHPWAPKRASMYSGAYNTAPDTPAVQPDGSPATGKRISPPNKRNLNGRKPSSASIVMIEHPQQRQLMPLQNK
REKGRDRVEKTDAAEFDPDTSHISKEAGSARSRTGSKMIMGCFRQKVLVRPSSASVPPESVVFELSEQ*
(SEQ ID NO: 159)

Name: Calcitonin receptor-like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013385192

GCTTACAAAACAAAAACAGATTTTGTGCTTCAATCACAACATAACGAAAGAAATATGAATGAAAAACAGAGGATTTATCATCATTGAAGGCTGACAAAACATTTGG
CACAATTTACATTTGCTCCGGCGATTTATTCGCGTCTAGAGACAAAAACACTCCGGAATTTGCTTGCAGTAAGAAAGTAATTTTAAATAGAGGTTACTGACG
TATGCAGCTGCTACTGCGGTTGCTACTAAAGCAAAAGTGAAGATATATCTTTGAATCTCAGCACAAATTCATTTCTCGTATTTTCTGCTAATGGAGTTGATGTC
ACCAAGAAATGCGTGAGTGTACACTCACAATAATGTTCTATTCCCAAGAAATTTCTCTCTGTAAGTAAGTGAAGGTGTGTACACTAACTCTGTTTGTATTAAGACAA
ATGCAATAAGAAATGCTTAAATTCGTCTACGTGTAACATGTAAGCCAGAAATTTTGTGATGTTACACGTTTCCACAAATTTGCCAAATCATCTGCTTCTGCCGCCAC
TAGCTGTTAGCTATTTATTTGATTCGGAACGAGGCTTGTCTTCTTACCTTTCCGACAAATGGAATGACATTTTCTGTTGATTGCTTTGAAATCGTACCTTTGACCTGC
CCTGAATTTGGACATCGTCTTTGATGTCGTCATGCAATGCCATTTGATTTGGGAATTTTCAAGCTTAAATTTCTGTTAAATTTCAACACCAACACTGAATTCGTTTCTTA
TTACCTTAACGCTGATGTTTATTTGAGGTAGCAGGTAGCACTTTCCGACGCGAGCTGCATTTTGAAGTGTGGAATTTCCGAAAAGGGAATTTAAATGAACGCCAGAG
GAGCAGCGGGGCGCACTGAGTTGGCAGCCGACACCGCGGTCAATCAACAGGAGTCTTTCAGGACATCCACCAACCGACTACCATCTGCGCTTCTATGACACCATGGTAC
GAACATGTCGACGCGCATTTATTCGCGCAGCGCATCGATGATTTCCACACAAAGTGAGTAGAACGAGGAGCTGAGCTTAGGCTGGGCACATAGCCCTGGGTATGCTGC
ATATTAATTAAGTTTAAAGTGAATTTACATAATGATTTGCCAGCCGGGGGGCCATAATGAAATCATGTGTGTAATTTGCCACGCGGCTGTTGGCAAAAGGCTAACGAA
GTGTGCTTAATGTGGATGGGCCCCCTTTCCGATTAAACTGAACAAATACTCGCATAAATCCATAAAGCAGATGTTAAAGGTGGAGATGAACATTTGAAAAATATATTGAGT
TCAATCTAACCAACCATGTTGTTTATTCGTTTCTTCTCTACATAAGCTCTAATTACTTTAAGAAATTAATCATACCAATTTTGTGCTCAAGTGAAGTGGTTTAAATA
TTTATCCCAATTTTGTCTTCTCCCAACAGCTACAATACTTTACGGTTACTTCTGCTGATTTGCTGTATCTTGGGAACCATTTGCAATACCTTAATATCATAG
TGCTGACCCGACGGGAGATGCGCTCCGCCACAAATGCCATACCTCACGGTCTGGCTGTCGCGCTGCTGATGCTGGAATACATACCTTATACGGTGCACGACTATAT
CCTCAGTGAAGGCTGCCGCGAGAGGAGCAGCTCAGCTACAGCTGGCGTCTTCAAGTTTCACTCGGTATTTCCCGAGTGTGCACACCATCTCCATTTGGCTAACG

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GTGACGCTGGCAGTTTGGCGGTACATAGCGGTAAAGCTATCCGAAAGGAATCGCATCTGGTGTGGAATGCGAACTACTCTGATACCATAGCCACGGCCTATGTGTCTGTG
TCC TGGTAGTGTACCTTGGCTGTACCTAGTCACAGCCATTCCGAAGTTCTTAGAGACTTTGGATCCATGGCGAAGACGATTGCCCTAGTGCATTAGTCAATACATCTT
GGACTACAATCGCAGGATGAGGTGACCTGACGGTCAAGTGTGCGAGTACACGCCAGATGTTTCTGGGCGATACCAAGTGATTGCCCAATGGAACTGCGATTAGCTTGTGTA
AGCTTACACCAAGTGATACCCCTAACCACATTAAAGCACTGGAGTAACCACATCTCTCGTCGTTGGGTGAGCGCAATGTGACTGTCTATAAGCTGTATCACAGCGCAC TGGCGC
TGGCGTATCGCGAGTTCAGGAATGCGACCTCTCTTATATACAGTGTCTGTATCAAGCTGATACCTGCTTCCGACTGACCACTTGTGCTGTGTGGCGCTCATGGTGTCTGTG
GGAGGCGCAAAAGAGGAGGAGAGATCTGCCCTGCTATGACGCAACGATATGACGCAATTGTGTAATGAAAGGTGGTGATTCCGACGCAACCAAGAGCTGTAAACTGTGTG
GAGAAGAGAGAGACACCGGTACGCCACAGGAGTGTCTTGGCGGTACTGTGCTCTTCTGGTACCAGGATTCCACAGGGCATTATGGGTCTGTGTAATGTGCTCTGG
CGATGCTTCTTCTGCAATGTACTTAAAGCTGAGTAAGTATTTAGGATTAAAGACCACTCTCCGCTGTCTGTAACCGTAATTGCTTTATTGGCTATGTGTAATGTG
ACTGAATTTATGTTGGTGTGTTAGGTGGGCCGATTTGGCACATTCAATCATACGCACGCTGGTGCCATTAATGCACCAATTTGAGGCTCAATGAGGCCGCTGG
CAATGACGTGTCAATAAGTGATCATTGGTTTATCTGCTACACCCAAATAAAGTAGTCTTACATTTAAAACCTTCGCAGATACATTAAATGAGAGATTTTTTCTTTAA
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ACTTTATGAAAGCCCTGCCCATAGATGGCGGTGTGGTGGAAACCCGACATTTATGGCATTTCGATTTTTCGTTTCAGGTGACCTTATGGACATCTGGCGCTTATTAATTC
AGCATCAACTCTATCTGTACTGTTCTGATGAGCGCCGATCTCGGAGACGTTTCGCGTCCCTCTCGTCCGCGCTGGCTGGACAAATGGCTGCCGCTGTGCGACAGCAGCG
CGCAAGGCGAGGTGGCGGAAGTGGCGGCTTGGCGGCTACGGCGGATATGACGCGAGCGGTGCTGCAACAGGATGCCGTAGCAAGACATGGCCATCGATCTCGGCTG
GACGACCCCAAGTGACAAATGTGTAGCAGGAGACGACGCGCGCGCGGCGATGTGACGCGAGCGCGGAGGACGCTGATCGGTGGCTTGGCCCTGGCAGCCACTGATGTT
GTGATGTGTCGCCCTGCCACCGCTGCTGTTTTCATTAACGACATACGCTGGTGTGAGAGACTTATTTGACGCGCAGTCCAGGGGGGACATGCGATATCCAGTGGCCAGC
ATCGAAGCGCGCGCAGTGGGAGTGGACCAAGTGCATCTGGCCACGACGAGGATGGCTGAGAAGAGCTGCGTATACCAAGAGCGTAGAGAAACGAGGACATCTCTCGAACAGGA
CATAGAGCTGGGCAAGAGCTCCATCAACAGGCGCAGCAGTGTCTTCTTATTAATGGTATAGTAAGCAGCTCCGATGAGGTTAAAGCCAGGCGGTTTGTGTCAGTGAACACGCG
CCGAGCTCAGCGGACGAGGATGTGGAAGACGCCATTGACGCCCTGCGCTGTGAGACGACCCCTCACTTCTATTGTTAGCGCCCTGTAAATATTCAAGTACCCACATTAAT
TTAACTACCGTGACAGACCAATTTATGAGACAAATGTTATGGCGTGATTCATATTTTCCGCGAGTGTGTGTTGGGCAATTCAGAGCCCTTTCGCTGAGACTTTGGC
TTTGGCTGGAATAGGCTGATTTTCTTGAAGGACTCTCTGGAAGCAGCTTCCGACACATTTTGTGACACATTTGATAAGTGATATCTGACTAAGTTTCCCCCTTAAT
GTTGTGTAAGTGTCAAGCGCGCTCTAGATATAAAGCAATAGTCTATGCTTAAC TATTAATAGATATATTGATAAAATTCGGAAGCTAAGAGATTAAATTTCTTT
TATATTGAATAAATGTTTTTTCGAAGTAAACTTGATTCAATTCACTTATTGTAAGTGTATTGCGACACTTTGTTTTAATTAACCGGAATTATATTGACTCACTTTGATTAACG
AATGCTTTGTAATGCAT
(SEQ ID NO: 160)

Exon: 1001..1062
Exon: 1492..2093
Exon: 2172..2612
Exon: 3102..3385
Start ATG: 1001

Transcript No. : CT33298

ATGGTCACGAACATGTCGCAGCCGCATTTTGGCGCACCGGCATCGATGATTTCCACACAACATACAATACTTTACGGTTACTTCTCGTGATTGTGCTGATTCCTGGGAA
CCATTGCGAATACCCTAAATATCATAGTGCTGACCCGACGGGAGATGCGCTCCCCACAATAAGTCCATCACTACCGGGTCTGGCTGTCGGCCGATCTGGCTGTGATGCTGGTAAT
CATACACCTATACGGTGCCAGCACTATCTGCTAGTGAAGGCTGCCCGGAGAGGAGCAGCTACGTTACAGCTGGGGCTGCTTTCACTCAAGTTTCACTCGGTATTTCCCCAGGTG
CTGCACCCATCTCCATTTTGGCTAATCGGTGACCGCTGGCAGTTTGGCGGTTACATAGCGGTAAGCTATCCGCAAAGGAATCGCATCTGGTGTGGAATGCGAACTACTCTGTATCA
CCATAGCCACCGGCTATGTTGCTGTGTCTCTGTTAGTGTCACTTGGCTGTACCTAGTCAACGCCATTGCGCAAGTTCTCTAGAGACCTTTTGATGCCCAATGGCAAGACGATTGCC
CTACGTGCGATTGAGTCAATAACATTCTGGCATACAATCGGCAGGATGAGGTTGACCATGCGAGGTCATGTCAGTACAACCGCCAGATGTTTCTTGGCGGATACCAATACCA
TCTCTGTGCTGGGTGAGCGCAATGTGACTGTCTATAAGCTGTATCACGCCGATCGGGCTGCGTGATCGGCAGTACGAAGATGGCAGCTTCTTATATACAGTGTCTCTGA
TCAAGCTGATACCTGCTTCGCACTGACCATTCTGTCTGTGCGGCTCATCGGTGCTCTGTTGGAGGCCAAAAGGAGGAGGAGATCTCTGGCCTGTCTGACGACCAACGATAT
GCAGCCAAATTTGCAATGGAAAGGTGGTGTATCCGACGCCAACCCAGAGCTGTAAACTGCTGGAGAGAGGAGAAGCAGACCGATCGCCACCACGAGGATGCTTCTGCGGCTACTG
CTGCTCTTCTGTGTCACCGAGTTTCCACAGGGCATATGCGTGTGCTGATGTGCTGTCGTCGAGGATGCTCTTTCTGCAATGTTTACCAAAGCTGAGTGACCTTATGGAACA
TCTTGGCGCTTATTATTTCTGAGCATCAACTTCTGTAAGTGTTCGATGAGCGCCAGTTCGGGAGCAGCTTCGCGCTCCTCTTCCGTCGCGCTGGCTGGACAATAGGCT
GCCGCTGTCGCAGCAGACGGCGAAGGGAGGGTGGCGGAAGTGGCGGCTGGCGGCTACGGCGGATATGGACGGCAGCGGTTGCTGCACACGGATGCCGTTAGCAAGAGC
ATGGCCATCGATCTCGGCTGACGACCCAAAGTGACAAATGTGTAG
(SEQ ID NO: 161)

Start ATG: 1

NVNTMSQPHYCGTGIDDFHTNYKYFHGYFSLIVCILGTIANTLNIIVLTRRMRSPTNAILTAKFLAVDLAVMLEYIPYTVHDIYLSVRLPREEQLSYSWACFIKFHSVFPQV
 LSTISWLTVTVAWRYIAVSPYQQRNIRCMGRTTLITATYFVVCVLVSPWLYLVLTALGLEFLTDANGTKIASVPLSQYILIDYNRQDEVTVMQMSSTPDVSWAIPITV
 SSGISGERINVTYKLYHSALALRDRQFRNATFLIYSWLKIPFCALTILSVRLGLTAEAKRRRKILACHAANDMQPIVNGKVPIITQPKSCKLLEKEQTDRTRTMLLAVT
 LLFLVTEFPQGINGLLVLLGDAAFFLQCYLKLSDLMDILALINSSINFILYCSMSRQFRSTFALLFRPRWLDKWPLSQHQDGEGRVGGSGGLGGYGYGRQRLHTDAVSKS
 MAIDLGLTTQVTNV*
 (SEQ ID NO: 162)

Name: TRH receptor-like
Classification: G protein linked receptor

Celera Sequence No. : 142000013384809

ATATACTCTTATGAAAAATAAAAAATGGGCTGATTAAAATGATTGGTATTATGGAATATTATTATGAAAAATATGAAATTTACATTTTAAAAAGTGTTATTCGCGTTATTA
CTCAGTTTGTAAATAGCGGGCACTCAATTGTGCATAGTTTGAAGGTAAAGTGCCTAAATAATGGCCGAAAGCTGTAACCGCGAATATATATATGATGTACATATACAGATAAA
AATAAAGTGTGGCCCTTGGCTTGGATAAAAAAATCGTCCGCGGATATCCATATTATCGCTTTAGAAAATTCGATATCATCGCATAGGTGTTCCCTTTCAAAAAGTC
TCGTCGGAATTTGGTCGTTTGGACTACTTTTATCTGTGTGATCCCGGACTTTGACTGCTTGGACGGAGTGGGATGTGTAATGAACACTAAAACGCTTGAAACTTGAGAAATTA
GGGAAGTAGACGGGTAGCGTCAGTTGTGTTGAGAAGATTAAACAAAGCCCAAAACTGCACAAAACATGCAAGGCCATTTTATTTATTGTCTATCCAGTTTCGATCGATATGGC
AACAACTTTAAGCCGATGCCCATCTCAATGCGCTCAATAAATGCGCACTAATCAGCGGGGCCCCAAGTTAATCTGGCCCTCTCATCGAGTCAGCGCTCTCTTCGAAGAAT
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TATTAATCCAAGTGAATTATAATATTTATTTTGGCACCTTGTAATGTGCAAGTGATTAGAAATGTGGTCTTCTTAAAGATATAAATGTACATATAATATGCTAGAGTA
CTGAGGCACTATTATTTATATATCTCCCGGTGTCGACAGACTAAATGATTCATAATGAATATGAATCCAGAGTTCACAGTAGGCAATAATCTCAGTTCCCTGTAAAGTG
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CATCGAAGCTTGATACCGGAGCATTAATAATCTCAGATCTGTCTGACTTGATTCCCTGTAAAAAATAAATGGGCAATTAATAATATTATAGCTTCTATTGCAATATTTC
ACTTACCAATTAATGGCCACCAGGATTGACGTGCAAAATCTTGCCAGAGGACGAATAAACACAGCTATGGCCAGCGCCAGAAATGCAATTCACATCCAGAATCAATGA
TGAAGAATAGCTCTCTCACTCTGGCATTATACAGCACTATTCTCGGATCGAGACGCAAAACAGCCCACTATGTAAGAGGCTAAGAGGATCGAAGATATAAA
AAGATAAATAGGCTCTTCAATAAAAAACAGTTTATAGCAATCTTATCGCTCTTGACAGTTTTCAGTTGAGTGCCTTATGCTAGATATCTGGTGGCAGTCTGATAGTGG

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AAAAGTACGCTTGTAGACAGCATACAATGCTGAATATCAGGAGGAGTGCATGGGTCGCTAGTAGTAGATCAAAATACCCAGGTGCGAACTGAAATGCATAGATAATTCG
AAATTAGACTTTTGTGTACTTTCTTATTCACCTACCATCGAACCAGCAGTAATCTTCTCCCATGCTGGCTTGAAGTGCTTGGGCAGCTTGGAGTCTGGGCTAAGGAGAC
CAAAAACCGAAGACCAATCACCGCCAGAACTGCCAGAGCAAGGACAAATCGACGGACGGAGCTCCTGACAGCCTTTGTTTAAATTTGAGCAGAAAGTCAAGACTGCAAAATG
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TCTGCAAGCCACATGCGAGAGCTTCTATGGGATTCTCAAGGCCAAGTAGACCAACAGGGAGTAACCAACGATCATGGACAGCAAGTAGTAGATGATTAAGTGGCCATAGTGG
CTTTTACGGGCATCTCTACAGATCCAAGTAGGGAGAGATGAAAATGTTTATAATGATGGCTATAATACTGCCTAGAAAGGAACAAGAGTTAGAATCTCTTCGAAAGTATT
CAAACAACACGACTCACAATGGCGTAAATCAAACCCGGTACCCCGTTTGAAGCGTTTCGAGTTGAGCGGAGTGAGCTCCACTGCTCCGGATTGTGCTCCAACGGGCTA
AAGCAATCTCATCCGTTGACCAAGAGCTTGTCCGCGCAGAGTTCCATTCTCGAAGAGGTCCTTGGCAGAAAGTTATCGTGCTTTTGTCCACGAATTTGTTCGGGC
AACCCAGCTCTGTGCGCACCACAAAACGATCGCGTATGCGTACTTGATCCACAGTCTGTTAGCGTAGGTGAGCTCCACGTGAGTGGACTCGCGGACTTGGTGTGGCATTGT
GCAGTTCCAGTGCTTGGCATCAAAGACGAGGTTCTCCGACACAGAACTGATGCAGGGCTTCAGCAGACAGACGCGCCGCGAGATGTTTCTGGCCCGGTACTCCACT
CCATCGATCACCTTGAGGAGTACTCCGCCATCAGATGCGGTGGGACCACTACTCCAGCGTATGAGTAGGAGCCATCTTATCCGAGCTCCATCCGTAATGTTGACCGTGT
GGCGTAGGCACAGGATGTTTATCTGGCGATCTCCACTGACTTTGCGCCAGGAGAGCCAAATAATTAAGAATTTAATGGCGAGACCATGGCTAACTGATCCAC
GCTGTGTGGATTTTGGTCGATTCTGTCGCGCTGACGTCACACGCTGTTACGCTCGATAAATAGTAACAAATATGAGAAGCGATCGCCGTTGCTTTGAGCAATGCA
CCGCACTCGCCGAGTCAGAAATCTGCACCAACAAATGGAAACCCAAATTTGACCAACGAAATGTGACGCGGGCTTAAAAACGGCTGCGATTGCGACACGACACCGACAAC
ACCAAAAGAACAAACATAAACAATAAAGTTTGGTGGAGGTCCTCTCCTAGCTAGTCTGCTTGCCTCTCTTAAACGACAAACCGGCAACCGCAACGGAGCTCCATGGCTGGC
CAAGAATAAAAAACAGACTAACTGATTAGCCGTTCCCAAACAGACGCGACCAAGCAGCTGGCAAAAGAGCGATTGGGAAGACTCACAACCCGTCGCCGCCCTCTCTC
TAGAAACCGCGCTGGAATCTCAACAGCTCGGAATTCGCATGCATACCTTAAGTTGTAAGCATATAATCCATTGAGCCATTTCCAGACTTATCTTGAGATAGATATAAA
AGTTAAGAAATATAGATTAGATTATTTCCATCTTATATAATTTATGGCTTTGGGACCGTAAAGAAATCCAATTTTAAACGCAATCCAATTTGCCCCAAACGAATTT
TACCACAGCTTTAGTTTTTAAATATTTAGGTGCATTTCCAGGCTATACAAAGTTTGTCTACATGTAAGGTGTGACCTGGCTTGATATCGTTTACGGTCTTCAAGAAATATTC
GAAAAGATTGTTGATTTTGAACAAAGATCCGCTGAGGAAAAACGACGTTCAATTTATCTTTGGCTTTATTAAGCTTGAATACAATTAATTTCTTGTATTGTGTTT
ACATTTTCAGAGATAATCAGATTAATGAGATTTTATGCAAGAGGATATACATCAACCAATCTCGTTTAAACAGGACAGATCTGATGTTACGATTTTACGATTTT
TGCTGGATGTTTACCTTGGACTTAGGCTAATGGAGAATTACAAATTTGATCTCTCAGCTGGGTTCCGAGAACCTTACAATTTTGTGTCATAAACAGTAGATATGCTCTGC
ATAAATTTCTTATACATTAATCAGGTTGTTCTGTCGATGCGATTAAAAATGAAAGAGAATTGGCGGGGGTGGGTGGGTAGGGTTTTCTGTTAGTTCTGGGGAT
AGGGACTTCTCTTATAGTAGTGTCTTTCATAAAGTTCTATCCGTATGTGGATGCTACTTCCGCAATTTA
(SEQ ID NO: 163)

Exon: 3214..2258
Exon: 2201..2010
Exon: 1950..1717
Exon: 1658..1502
Exon: 1442..1239
Exon: 1176..1097
Exon: 1035..1001
Start ATG: 2897 (Reverse strand: CAT)

Transcript No. : CT33414

TCGTTTAGAGAGAGGCAAGACGACTAGCTAGGAGAGACTCTCCACCAAACTTTATGTTTATGGTTGTTCTTTTGGTGTGTGCGTGTGCTGCGAATCGCAGCCGTTT
TTAGACCCGCGTCACATTCGTTGTCATTTTGGGTTTCCATTGTTGTTGGTGCAGATTCTGACTCGGCGAGTGTGCGTGCATTGCTCAAAGCGAAGCGGATCGCTCTCA
TATTTGTTTACTTAAATATCGAGCTGAACGCGTGTGACGTCAGGCGGACGAAGCAATCGACCAAAATCCAAACAGCGTGGATCAGTTAGCCATGGTCTCGCCATTGATAA
TTCTTTTAATTTATTTGGCTCTCCGTGGGCGCAAAGTCAGTGGAGATCGCCAGTATAAACCACTCCCTGTGCTTACGCCACACGGTCAACATTACGGATGGATGCGGATGAA
GGATGCTCTCTACTCATACGCTGGAGTAGTGGTCCCACCGCATCTGATGGCGGAGTACTCCTCAAGGTGATCGATGGAGTGGAGTACGGGCGCAAGAAACATCTGCGCGGC
TGCGTGTCTGCTGCTGAAGCCCTGCATCAGTTCTGTTGTCGGGAGAACCTCGTCTTGTATGCCAAGCACTGGAATGCAATGCCAACACCAAGTCCGCGAGTCCACTCAG
TGGAGCTCACCTACGCTAACAGGACTGTGGATCAAGTACGCATACCGGATCGTTTTGTGGTGGCAGAGCTGGGTTGCCGGAACAAATTCGTGGACAAAAGCACGATAA
CTTCTGGCAATGGGACCTCTTCGAGAATGGAATCTCGCGCGGACAACTCGTCTGTGGTCAACGGATGAGTATTGCTTTAGCCCGTTGGAGCACAAATCCGGAGCAGTGGGAG
CTCACTCCGCTCAACTCGGACCGCTTCAAACGGGGTACCGGTTTGGATTACGCCATTTCGAGTATTATAGCCATCATTATAAACATTTTCATACTCTCCCTACTTGGAT
CTGTGAGAGATGCCCGTAAAGCCACTATGGCCAGTTAATCATCTACTACTTGCTGTCCATGATCGTTGGTACTCCCTGTTGGTCTACTTGGCCTTGAAGAATCCCATGAA
GCTCTCGCATGTGGCTTGCAGAAACATAGGATTTCTGGCCTACTTCTGCATCATGCTCTCTCTGCTTCTGCGCCATTTCGAGTCTTGCATTTCTGCTCAAAATTTAAACAA
AAGGCTGTCCAGGAGCTCCGTCGATGTTCTCCCTGCTCTGGCAGTTCTGGCGGTGATGGTCTTCGGTTTGGTCTCCTTAGCCAGGACTCCAAGCTGCCCAAGCACT
TCAAGCCAGGATGGGAGAAGATTACTGCTGGTTCGATGTTCCGACCTGGGGTATTTTGTATCTACTACTACGACCCATTCGACTCTCTGATATTACAGATTGTATGCTG
TCTCAAGGCGCTCTTTCCATCTACGAACTGCCACAGATACCTAGTACATTTGGGACCTCAACTGAAAATCGTCAAGACGATTTCTATGCTTTAGCGCTTACATAGT
GGCGTGTGTTGGCTGCTGAGTCCGAGAAATAGTCTGTATATAATGCGCAGAGTGAAGAGCAATTTCTTCATCATTTGACTTCTGGAGTGGCAATTTGATTTGGGCTGGCC
TAGCTGGTTTTATTCTGCTCTTGGCAAGAAATTTGCAGCTCAAACTCTGGTGGGCCATTATGTGGAATCAAGTCAGACAGATCTGAGTATTATTAATGCTCGCGTATACAA
GTTTCGATGAGAAAGCGATCTAAATCTCAGACTCTCCCTACAGCGGACAGTGACATCACTTTA
(SEQ ID NO: 164)

Start ATG: 318 (Reverse strand: CAT)

MVSPILILLIWLIVGAKSVEIASINHPCHAYHTVNITDGLRMDGSYSYAGVVVPPHMAEYSFKVIDGVEYRAKKHLRGCVLLKPCISFCCPENLVDAKHWNCTMHPQ
VRETHVELTYANRTVDQVRI RDRFVVRTELGCNRKFDVKKHDFWQWDLFENGTLRRDNRLWSTDEYCFSPLEHNPEQWELTPLNCERFQTYGRVWVIAICSI IAI IINIF
ILSLGSGVRDARKSHYGQII IYLLSMIVGYSLLVYLAKNPMKLSHVACRNIGFLAYFCIMLSFVFLAICSLDFLLKFKQKAVRSSVRRLLSLAVLAVLIGRLVLSAQD
SKLPKHKFKPMGEDYCFWDRVTGILIIYYGPIALLLIFSIVCLLKFYSIYELPPDTQYILGTQLKIVKTHFYAFSAIYGVFAVWIREIVVYIMARVREHFFIIDFWSGI
CILGLAIAGFILLGKNLHVKSWWAINVESSQTDLSIINARVYKFDEKGDKSSDSPKPPVTSL*
(SEQ ID NO: 165)

Name: mth-like 9
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013384809

CGAGCCATTTCAGACTTATCTTGAGATAGATATAAAGTTAAGAATATTAGATTAGATTATTTCCATCCATTTTCTATAATTTATGCTTTTGGACCGTAAAGAAAATCC
AATTTTAAACCGCAATCCAAATGGCCCAACGAATTTACACAGCTTTAGTTTAAATTTAGGTGCAATTTCCAGGCTATACAAAGTTTGTCTACATGATAGGTGTGTGA
CCTGGTGTGATATCTGTTTCAAGGCTCTCAAGAATATTGAAAAGATTCTGTTGATTTTGAACAACAGATCCGCTGAGGAAAAACGACGCTTCAATTTATCTTTGGCTTTATTA
AGCTTGAAATACAATTAATCTTGTATTGTTTACATTAGTTTACAGATATGAGATATGAGAAATGGTACATTTATGCAAGAGGGTATACAAATCAACACCAATCT
CGTTTTTAAACAGGCACTGATGTTGATGTTTGTGTTGATTTTACCTTGCATTTAGGCTTAAAGGAAATACAAATTTGATCTCTCAGCTGGGTTCCGAGAACCTT
ACAATTTTGTCTGCTTAAACAGATAGTGGCTCTGCATAAATTTCTTATACATAATTTAGGCTTGTGCTGCTGTCATGAGCAATTTTAAAGTAAAGAAATTTGGGCGGGT
GGGGTGGGTAAAGGTTTTCTGTTAGTTCTGGGGATAGGACTTTCTCTTATAGTAGTGTCTTTCATAAAGTTCTATCCGTATGTTGATGCTACTTCCGCAATTTATAGAT

FIGURE SHEET 72

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Exon: 2646..2805
Exon: 3377..3577
Exon: 4022..4372
Exon: 4614..4820
Exon: 5539..5804
Exon: 5893..6112
Start ATG: 1001

Transcript No. : CT33559

ATGCAAGATCTGTCCATCTATCTGATAGAAAGCTGGGCTCTTTGGCGAGTTCCTCTGCCGCATGTACCAAGTTTGTCCACTCCCTGAGCTACACGGCCCTCCATTTTCATCTCTGG
TTGTCATCTGTATGCGAGCGCTATCTTTGCCATCGTACACCCCATTAACCTGCAAAACAGATCCTAATCGCGGCTCGTCTGAGGATGGTCATTGTACACAGTTTGGATTACATCGGC
AGTTTACTCCACGCCCAAGTTTGTTTTCAGCAAGACACCTTAAGAACATCCACACGCGAGGCGCGGAGGAGATATGTGCTCTGGCCGCTGAGATGTTCAACTCCAAC
TTGCTGGACATGATTAACTTTGTGCTTCTCACTGTAAGTTCGCGCTCTGGTATGACGGTGTTACAGCAAAATCGCCATGCCCTTTGTGGCGCAGTTCGCGCGGCTCACGC
CGCATGTGGTGACAGATCAGCAATCAGCAGCGCAGCAGCATCTCTGGCAGGACATTTGTCATGGGTATGCACAAAGCATGTACCATACCACTCCGCATCATCATACCAACCA
CACACAGCATCACCAAGCTGCAATCGCGGGCTTCGTGCGCAGGATGGTGGGAGTGGGTTTGGGTGGTGCTGGTGGGGCGTCTGGTGGCTTCGCTCGGCGCGCAGC
AGCACACGCTCTTGTCCCGCAACACGAGCAGTAATAATGAGAAGCGCGGCTGAGCATACGAGAGGCCAGTTGGCAGTCTGAAAACTTTACCCGTAGCTGTTACTCCCG
CACTTTCGCCGGGCCAAGTGGCACGGTGCTCGGAGCAGGTGTCAATTCACATTAAAGCCATCGCAGGACCATGCCACAAAGCTCGGCCACOGATGTGACACCTGGAGTGTGGA
GTGGTGGAGCGAAGTGAACACCGGCACAGGGCGAAACTGCAAGTTCGAGGCCAAGTTCCTCGTGGAGCGCATCGCTCCATTTGTCAGTGGCTCGCCGCAAGACGAGCTTCTAC
CACACCGGCACCGCCATACCCAGCGCGGCGGCAACGCTCATGTGGCGGTGGCAGCGGCGCGCGGAGCCACCCATGATGTCCCACTGCTCCAGCAACGCTCTCAT
GCGCCCGGCGCGGCGTGTCCGCATGCTGATAATATTCTGCTGACATTGCGCCCTGTCGAATCTGCCGTATCATGCCCGCAAATGTGGCAGTACTGGCCATTATCAGCCC
ACACGGACTGAGTGTAACAATATTTATGGTATATTGCACCGGCAATTCGCGGGAATTTGCATAGTCTCTCGTTGAGCGGGAATTAACATCTGCGAGTGGGCAGATAAGC
TCTAGATGGCTTCTGTGGAATCGAGGCTAGTAAACAATTAAGTCATCAATGGACAGAGCTGGAGCTGCAGCCAGCTTTGCCGCTGGCCTAA
(SEQ ID NO: 173)

Start ATG: 1

MQNLSIYLIESWVFGEFLCRMYPQVHLSYTA SIFILVVICMERYFAIVHPITCKQILTAARLRMVI VTVWITSVAVYSTPKFVFSKTIKNIHTQDQEQEEICVLDREMFSK
LLDMINVFLLVYMLLVMTVLYSKIAIALWRSSRGLTPHVQHQQHQQQPSCQDIMGMMN SMYHHHPHHHHHHHHQHQQLQSAASSAGVVGVLGGGGGGGGPGLSASGGS
STSLSRKQVYKPEKRGVSI TESQALAVKTL PVAVTPALSPGQVARKSEQVSIHKTATGPGCHLGHRCDTSPVSEWVSEVTTGTGRNCKCEAKVSLSEADRPVISA CRKTSFY
HHGHAAHHQRAGNASVGGGSGGAGAGATHMSSSSSNVLRRRGVVRMLIIFVLTFALCNLPYHARKMWQYWPFI SPHGLSVNNIILWYIAPAIGGNLHSPPLDGLKLTASGQIS
SRWLPAGIEASKQLSHQWTELELQPALPLA*
(SEQ ID NO: .174)

Name: Neuropeptide receptor-like
Classification: G_protein_linked_receptor

Celera Sequence No. : 142000013385217

[illegible]

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GATGTATAGGCATCAATACGGATATAAGCAGAAAATCTAAGATTAAACATTTATCAGCAAGTCTGCAGGTGATGACTTAGTAGTATACCATGATGGTGAAGATTGCGAGA
TATAGGACAAACCATGAAAATCTTAAAGATCAACGAAAGAAATCATAAACACAAAACCGAACAGATGATATACCGTTGATATAGGACACAGCGATGATCGAGTAAGGGAA
ATTGGAGTGAATACTAAAGATTACCAAAATCATTAATACCACCAATTGCGGAGATGCATGTGCACAAAATGGCAAATTAAGGGACATTGGAAACGACAGATAGGCAT
TTTGCCCAATCGATGACGGAACATGATTAATCTACATGCATCCCATTAAAGACTGATAGGAAGAAATTAACAAAACATGATTGGATCCACCAGCGGATTAATGGACCTTACAA
GATGCCAACCAAGAGATAGACGAACGCTACTACAAAGGTTGTGAATATCATTTTCTGGCCGACGGAATGGCGCAGGCTTTTCTTCAACAAAATCCATGGAAAATACAAAG
CTACGCCGCTCCAGTCTATTGGCTCTACACACTAGTTTTCCGTTTATATATTTTGTGTTGTCATTCTTTCCATGGCCTGGTTCGATTTCATCAAGATGACGCTTCCA
GAAAGGTACCATGATAAAGATGGCACAACCATTTATTAGTTTCACTCCCATCGGACACGAACATAATCCCAAGGCGGTATCCTTTGATCCTAGAAACAGCACAGAATCAT
GGAAAAGTACGCTGGTATAATGGCTCTTCTGGAAAAATACGGCGACTATGGACACAATCCCGCTTTTGGAAACGTGACTGCGAATGAAAAGTTTGGCTATCCAAAGCGCGGAA
CCGTGTGTTTTTCTTAAGTTAATCGGATTATTGGATTAAAGACCGAACCTACATCAATCCGATGAGCTTGTCAAAGCCAAGATCGATGAGGTTGAATTTACGGCTTAA
AAAGGTTATTGGAAAATACAACAACAGAGAGGGGTCATTGTAATCGCACTTGGATAACATGCGCTTCAGATAAAGATAAGAAATGTTTGTATTGAATTTTCATCCAGAGCGCGC
GATAAGAACGGAATACACCGATATTGAAGAGAAAATCGATATATTGCAAGCAGGGTAAAAAGTCATTTTCCGCCAAACGATGTGAATCGAATCGTGGCTCTTAAAGATC
AAAAATCTCAAAGCAATGGAGCTGTCCATATAAAATGCAAAATGTGGGCTCAGAAATATACATATAGGAAAGAGGGTTATGGCCAAGTTTCGTTTTTGTGTATTAGTCA
CAAAATGAAAACCGCAGAGGGGTGAAAAATTTACTCCGAACCGACTACTCATTATAAAATACATAAATAAATTTCTTAATAAAATGTGTTATATGATATATGATATGATTAT
TTATAACATCTATAATCCAGGTTCTTCAAGATCGCACAATTTGTCACCATGTTCTACGCAACCGTCTTCCATTATTGTTGTCATGTGCACAAATATCCAGTTGTGGCAT
ATTCGTGCTTCCGCTTGCCTTGTATCATCTTTAATATATGGAATAAGCATAGAGGTAACATTTAAATGGTAGTGCTCGTGCAAGAACTAATCAAGGGTTATATAATTTCTAGAGT
AATACAATCCTCGCATCCCGTTTGAATACGATCATGTTATTTGGTGTCTATCTGTCTAATATCTGTCTATCTTACTGGGCATCGACGAGCGCTTTGTGAGCCCGAGGAA
TATCCAAAGGTGCGAGTTACTTACCTTAAAGTTCTCTTCAATTTTATTAAGTTAAATTTGATATAAAATTTATCCATTAGATATGTCAAGCGCGGGCTTGGTTACTATCCAC
CGGTTTTACACTAGCATACGCTGCTATGTTTACGCAAGGCTGGGCTGTGCTATGTTTTACAAACAAAAGCAAAAACCTGACCCAAAGGTGGGTTAAATTTAAATTTACGCAAC
TATATATTAGTTGCTAATCATTTTGTGAATTTGCAGAAAAAGTGAACCTTGAAGCTATACACCATGTTTCCGGGCTATATCAATAGATTTAGTGATATTACTCTCAT
GGCAGATCTTTGATCCGCTGACGCTTATCTCGAAACATTTCCCACTCGAAGATCCAGTATCTACTATGATGATATTAATAACGTCAGAGCTTGAGCATTTGTGAAGTCA
ACGCAACTCTCAAGCAATGGAGTGTGTCATATGATATGATATGATATTTATTTATTTTGTGAGGCTCTGTATGATTTTGTGCTTCCATAGCATGCTGCTGATATTTGTGCGAAAGGT
TTTGGCCTCTTTTGGCGTACGAGACGCGCTCCATTAAAGTGAACAGATCAACGATTCGCGTTATGTTGGCATGAGCATCTATAAGCTGGTCTGCTTTTGGCTGATAACAG
CTCCGCTGGGCTGCTGATGCTACGCAAGGACGCGCTCTTTGGCTCTGTTGCTCTAGCTGTGATATCTGTTGTTTCCATAGCATGCTGCTGATATTTGTGCGAAAGGT
CAGCCACTACACTGAACATAAAAGAAAGTAGCACATTTGTAATGATCTCTATTTCGCACAGGCTATTGAGGTTATACGTCATCCCAAGGATAAGGCCGAATCGAAATACAATCCC
GATTGAGCCATATCGAAAGAGGACGAAAGACGCTATCAGAACTTGTACCAGAAACGAGGAATGCAACGATTAAATACACAGGTATGCTGAATTTAGGTAGATTATCTTA
ATAAAATCGTATTTTAAATTTTGTTTAAAGATTAAATTCACCTGTTGCTTGAACGTTAATCTAAAGTATTTCGGTCTTATCATATTTGAACAAAAGTTAAGGATTGG
AATTTCCAGCGGCCAGTGAAATTCATTACGATGCGCGTTTCCACTTCTCTAAATAAAGTCCATTTAGCGAGCTGCAATATGAAGTATTATCAACCTCTTATAAATCAA
ATTTTAATTTGATAATTAATTTACCTTGACAAACATGAGTGAACACCAACACCGCCATGCTCGGCTGCACAAATTTTACTAGATTATCATTTATCCGATCAAAAGTTG
AGAACCTTTTATTTATATGTTATCTTAGGAGTCCCAGCTGTTCCAGAAATTTTCCCAACCGTTCCAGTTTGTATAATTCAGTTTCACTGCTCCACTTGAAGTCTATCAAA
TGAGCGTATACGTGTAGCGCCAGGACCATGCTTCAATTTGAATGAAAAGTTGCTGTGGTTGTTCTCTAGTCTATTAGGTAAAGCTTTCGAAGACCGATGAAAATTTGCGCATTC
GGATCGCCAAATCCATCTTTGTAACATAAACCCAGATTGGTTAAATTTAACACTTCCATCTAGTCTGCTTTGAATGACAAATCAATCTCTTGGCGTCTGTTTGCATGGGTCA
CTTGAAGAGGTTCTTCTTCTGCGCAACATTAGCTGGTGAATTTCCATTGGAATGCGCATAGGACACGCTGTGTTGTTGAACCTTGGCCCACTTGAACAGCAAAAGTCCATAAA
TAGTACAAAAGACAGATTTCTTGTGATCTCGACATGCTGACTGTTAGAAGATCGGTGGCTTACTGAAATCTTACCATTTAAAGGGGAACATTGGGACCGAGTTTGTGG
TTTTGCTACCTTTGACCGCATGCTTGTATGATATATTTCTATTTAGGTCGGCATTTAAGCAATGGATTCCAAAAAGTGAATGGAGACTATTTAAAGATAGAACTACA
TTTTTATGTTTAAATGTTTGTGTTTGTGTTTATTTATTTGTTTATAAATGATTGATTTATTAATTTTCCGCTTGCAGAAAGGAGAAAGATTCCAGTCTTGCAGAC
CGTCTGGTGGAGCGGGGACGCCAAGGGCACAGAACTGAATGGTGTGCAACAGGTGTGCTCCCGCGCCCTTGCAACAACTTCGCAAGCCGCTTCCCTCATCAACTCATCAG
CACATGCCACGCCCGCAGCCACACTCGCAATCACACAAGGTGAGTAGAGTACCTGGTGCATGGAATATGCATCCCAATGCCCTTAGCCAGTTGCATGTGGTAAATCGATA
CTAGTTACGTTGGTTAGCTTACGTTCTTAATCGTTATGTTCTCGATTTTAACTAGAAATCTTAGCTCTATTCGTGCGCATTGACACAATGATTAGGAACAACAAATATA
ATGAGTTCAAGGCTATTGCTTATGCTAACGCTATGTAACTCGATTTTTCGCAAAAAACCAAGGCAATTTATCAGTTTCTGGAATGCTATTTGCTTTGTGTGCTATC
ATATGTATCTTAAGCACTTTTGCAGTTTGTAGCTTATTTATATCTTCAGCACTTTAGCACTTTTAAATAAATATCACTGCTTTCGACCCCTTTTTCACATTTTAGCAAG
ATTATTCAAAACATACTAATACGTAATATATTATCTTACAGATACATGAGCATCGTACATGAAAAAATGTCATTTTCCCTTTGCGTGTAGGTGAATTTATACAAAAGA
TATGAAATGCAAAATATGATGCTTAACGGAAGGATGGATTAAACAAACAGTACTTTAAAGAACTTTGTATCAACTTTAAGCTGAGCTAAATAGTGAATTTGGTAGCAGCAAT
TTTGTAGAATAATGAAATATGATTGACCTTAAACGTTTGTAGCATTAAATTTGAAAACGAAGTAAATTTAACTTAAAGAAAACAAAACAAAATTCGACACTAGCATCT
AACTAAGCGAATTTGTAGCTTTAAAGTTGCAACACAGAAACCAAAAAAATTCAGCAATCAAGAGAAATTCATTACGAAAGATCGTGTAAAGTTAGTAATTTTAA
TGCAGGTAGAGCACTGTAGCCGAGGCTGCCAAAATGTGGAATCTAATTTGTGCTCTGCTACGGGGCTCGAGTCCGGCTCTTTCGGATCGCAACGATTTCCCACTC
TAGTTAATTATATCTGCGTAGATGAAAATTCCTCAAAAG
(SEQ ID NO: 175)

Exon: 1001..1090
Exon: 1165..1208
Exon: 1810..2052
Exon: 3783..3856
Exon: 5748..6002
Exon: 10046..10180
Exon: 10271..10441
Exon: 10503..10719
Exon: 10812..11284
Exon: 11344..11493
Exon: 11565..14044
Exon: 14446..14569
Exon: 14902..15028
Exon: 15086..15342
Exon: 15403..15540
Exon: 16659..16847
Start ATG: 1026

Transcript No. : CT35221

TCATGGAGTTCCAGTCTGCGATACATGTGGAAACACACGAACACAGCAGGAGGCTCTCGCGTCTCATCCCCGAATACAATGCGAAAAAATGAAATACCTGCAAAAAGAA
ATACGCGTTAAATGGAATTACGACCCCCCTCCATCCAAACACCCCCCTACGACCCTGCCCCAAGCAAAAGAGGAGCGGAAAGAAAGAGCGCTCTTATGAAACAATTGAAGCACC
GGCCAAAGGAGCAAGGAGCAGACAAAGACAGAGACAAAACATCGGCAAGGATAAAACGAGGCGCGAGAAAGAGAGGAGCAGTAGTAGCAAAAGCAGGAACAAACAACAA
CAACAACCGCAACATCGCATTTTAAACGCGCAATACGCAAAAAAATGCTGCGCAGGAGTTTGAAGCAATGCCCAGGGGATGCGGACAGATGAAGGGGCGGGGGCTGG
CGGGATATGACAAGTGATGGTGTGCTTACGTTTGGATATTTTGGTTGTTTAACTGCTGCTGCGCCACCTGCAAGGGGGCTGGCCGGGAGGCGCGATGAACCTGCACATCG
GCGGCATCTTTCCGATAGCCGCGCAAGGAGGATGGCAGGGCGGCCAGGCGTGTATGCTGCGCAAGACTGGCGTTGGATGATGTCAACAAGCAGCCAAATCTGCTGCCGGG
CTTCAAGCTCATCTGACAGCAACGACAGTGTGTGAGCCCGGTTTGGCGCGCAGCGTGTATGATATCTGCTCTATAATAACCGCAAAAGCTGATGCTGTTGGCAGGA
TGACGACATGCTGACCACTGTAGCCGAGGCTGCCAAAATGTGGAATCTAATTTGTGCTCTGCTACGGGGCTCGAGTCCGGCTCTTTCGGATCGCAACGATTTCCCACTC
TATTCGCAACCATCCATCGGCCAGGTGCACAATCAACGCGCATCAAGCTGATGAAGAAATTCGGCTGGTCCGGGTGGCCATTCTGCAGCAGCGGAGGAGGCTCTTAT

FIGURE SHEET 78

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[illegible]

Exon: 5938..5903
Exon: 5607..5368
Exon: 5307..5249
Exon: 5009..4890
Exon: 4827..4747
Exon: 4669..4321
Exon: 4260..4192
Exon: 4130..3729
Exon: 3586..3392
Exon: 2213..2086
Exon: 2027..1916
Exon: 1543..1523
Exon: 1198..1001
Start ATG: 5938 (Reverse strand: CAT)

Transcript No. : CT35779

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Start ATG: 1 (Reverse strand: CAT)

[illegible]

Name: Metabotropic Glutamate-like RECEPTOR
Classification: G protein linked receptor

Celera Sequence No. : 142000013384249

CTCGTGACCTTTGAACCCGACGCTGGCACTTTGAAACGGAAACGGAAACACACACTCACACACACGTAATTTGGTTAGTCTGATATCGATGCACATATATGTTGGCTAATTTA
GACGGCACTTTGGTTTGTACCATCGCCAGGGTCTCATATATATGTTTGGACTATATATATCTGCTGATTTAGTTAGTTATACGATTTAGGGGTTTACTTTTGGATATTTTG
AGCATTTTCAATTCATGCATTGCTGTGTAAATACCAATTTGGGTTTGTAGCTTTGCTTTAGCTGCGTGACTTTTGTGTATCTCTATAAATTTATCAATATTCCTCCGATTT
TAATTCGCTTTGGGGGTGTGGTTGTTTATTCGACGAGACTTCTCACGTAGTTGTCTATAAATTCGCGTCTCTTAATTGGCACAGTTTATGTTTACGTTTAGCTCGTCGATTTT
GCGAATCCCTATGTGACACGCTAGCTCAGTAGTTGTCAGTAGTATGCCATTTTGAATAAATTTTGTGGTTATTTCTTTTCGTGACGAAAAGTCGTTATCGGGCTTACAGATTT
TCGATTTGGCGTACGTACATACATATACATATTCATATACATTTACGAATAGAGTTTACACAGTCTGTAATTTAGTCGTACAAAAATCTCAGTTATTCACACAGATA
TAAACGCACATAAAATCATAAAATAAATAAGTTATCCTATACAAAGTCCCGATCTGTGCTCCAAGGAACCCGGCAAACTCGAATTTGAGATTAAATGTTTTAAAAACGAAAAACA
AGATGAAAAAATGTGAAGAAAATGAAATGGCCAGCTGCACAGATTTGAATATGCAACATTCAGAGTAATACGCCGACGCAATTTGCCGGCATATTAATTTTACAAATTAAT
GGAACCACTTTAATGGCTTAATATGATTGACCCATCCGGCTGGGACGATTGATTCGATTCCGATCTTACACAGTCCGACCAATTTGCACATTTGACCAATTTACACAT
GACTCGAGGGCGTCGAGGCGGGCGGCAATTTGTTGTTCTGTTCATTGTTGATGTAATCCATGGGTATTTCAATTCGGATCTCCTGCACATCTGTCGCCCGCCAGCAATCGAC
ATTGGATTGTCCTTTGCCGCGATCTTCCGCCGATCTTCCGCTATCTTGGATCCTTCGACTCTCGCTCCGATCTTTGGCGGACGTGCCAATTTGCCCGCACGGCGGATCTCGCCCGCCGCG
AGCGCGTCTGCTGCTGCGGACGATGGGGGTGACGACGGCCCTACCTCGTATTATAGTTGTCCACCATCTGACGCTGGCGGGCATCGAGGGCGTTGTGCAGTTAAGGTAGC
TCTCCGAGTAGACGCTCTTCCGCCGCTGGTCTGAGCGAGGCTGGAGCGCCGATGTGCATGTCCAAGGCCAGTGGCGATCTTCCACCAATTCAGAACAGGCAGAGCAG
CGTGTTCGCGGAACGCCCTCCGGGAAGTCGCGGTTTGAAGTAGCCGATAGACAGCGGTTTACGCTGAGTTTGAAGTAGCCCGATCCAGAAAGAGGATTGAGACGACTTATGTCGCGG
ACTTGGCACTCTCCTCGAGGTGATGGAGAGGTGTGACTCTAAATTAAGACATAAATTAATTTAGCTTACACTATTACTTTAACTTAATTTAAGTCAATTTAAATTTAGTTG
CACAGGTAATGCAATTTTATGTACTTTTTTTTCCCGGCAATTCGATAATATCTATTTTAAATAACAAAAGCAAAATATAAAGAATCAATATTTTGTTCATAAA
AGTTAAACCTCAAAAGCATCTATATTGATCACTTTGATCCACCTTTGAAATATTTTGTCTGTGAGCAATCAAGATGAGATATATATATCCCAATAA
TTCGACATGTCATGAGTGGCGAGGCTTTTGCATAAATGAGATTAATCTCGGGTGGTGTGAAACAAACAAACCGCGCAACATGGCGCAGCGGCGGTATGAGTAATATA
ATTAATGATGCTTGACATTAGATAAAATATGCTTGGAAATCCCATGTGACGGGGAATGGCGCTCCAATCTCTCAACCAAAACGTTCAACCTTCGACGGCTGGTTAGTACTCACCA
GGTGAAGAAAGGCGACGCGACAGGAGTAGAGGTGCCCATGATGATGCCAGCGTGGCTGGCGGCTTGTGCTCCGGTCTATTGTTAGTTAAGTGTGCTGCTTGTAGTGGGGGTG
GCTCTGCTGCACTTCTGTGACGGTCAATGTTTTCGACGAGCCGATCGCTGACGCGCTCGCTCGCGTAGGAGATAGATGGAGATTTGATAAAAGTCCAGGCTGGAGAT
GGGCACTGAACTCAATGCATCGCACTGCATCTGTCACGAGTCTGTTGTAAGAACAACTAGTGATCTTTCACGCTCACTACTCTGTTGCTATTCAGAAAGTAATAGACT
TCTCTGCTCGGTAAATTTATATTTTGAAGTATTATTTCCACTTAAACCCCATATATATATTTCAACCAAAACATTTCTTTACGCCAAACTAGGTAAGGCTTGTGTTTAC
ACCCAACTTTGACTCCAACTCACTGATGGCTCGATGATGGCTGGTGGCTACGATGGCATTCGCTGGCGGATCATCGCTGCTCTCTCGCGAATGGCTTCCCGGAA
GATGGCCAGGTAGGTGAATATATGTTGCGAGGTAATCCAGAACGATATGGAGCTCGAGATGACGGCGTAGTACTTGTTCACCAAGAACAGGACATGCGTCGGATCTGTG
ATGACGAACCTGCTGGTGTCTCGGCGTGGTGTACAGCGCATGAAGTAGGGCAAGGAGCAGTGGCCGGCATATGATGTGTTTAGCAGCATATGCGCCACCAACGCGTT
TCGTCACTGCTAATCTGTTATGTTAGGCGGCTTAACATAGCATGATGATCTGTGAAGTGGGCAAAACGCGAGGTATTGGCGCCTATATAGCATAGTTCTGCTGTGGCAT
CAACTTACCTATCCACAGATATGCAGCATAAATGCAAAATCTCGCTGTTGAGAAGTAGACATCGAGGCTGTTTCCAATAATGCACAGGAAGGGGCTGAAGTCCACCTGAA
AGGTTCCAAATGAATGGAGCATATGATTCGCGATGCTTACCCAATACAGGAGGCAATAAACTTTGGCCTTTACTTTCTAACGCTATTGATCTGATTATCGAGTCCCAATG
TCGATATATATATTTTGTGTGACTCTTTCTGTTTTTGGCTTACCATAAAATGCCAGCGAAATCTTTTAGAGTCATCGCTTTAAAACTGAGTTCTCGTTTGTAGTCT
TGCAAAAAATTTAGTTTTTACGATTAATTCCTTTGTGACAACACAGTTATTTACGTACTCTGATTAAAACTATTGATGTTGGGTAAATTTGCGCTAGTTCTAATATTTA
CTACCATTTAAAAACAGATATATAGGCCATAAAAACTACCTAGATATGCGCAATAAGGTTGCTCTAAATTTAAACGGCAATTTTACTGCCAATTCCTGAATGTAGGTCATA
AAAACTCAAACCTGACTAGACACACTCTCTTTTTCGCTATTTCTGTTATCAGCGCGCTGATTTATGTGTTTATCGATTTCATCTTAGTTTCTGGTGGCAGGTGACATGTC
CCCTGAACGCCATTCTGGAGGGCCAAATTAAGAGCTTAGATGCATCTGTGAGACAGAAATAGGCCATTAGGCGCGAGGTGAAAGGTGGCAAGGAAGCGAGGCGCACTCT
GTTCGCTGCATGAAAACTTATAAACTTCGCTGCTGCTGTTCGCTCTGCTGCTTCCGCTTGGTGGCCACTTCGCGATTCGTTTGGGCGATGGGCGGAT
CCCCGCCTTTGGGGCGAGGACGAGTGAATTTATATCGGGCGCCCTTGGCATGAGTTTCAAAACATTTTGCATATCCAAACAGGCTGTTTTTGTCTGCTGGTTT
TTGGTGCTTTACACTACATTTGCCATTTGGGCGGGCGGCCATCGGTTTGAACATTAGCCATGAAATGGCATTTCTCGTCTGCAATTTATGCAACTGGCAACTGTGAAG
GATAAATGCTAATGAATGCCAAGTGGCGGCGAGTTGGCTGTGGCCAGAACAAACCGATGCCAAATTTATATGCAACTCGCGCTGGCCAAAGTTTTTC
(SEQ ID NO: 184)

Exon: 3243..3209

TTCCCGTGC CGCGACTGTTGCGCGCTTGAGTGCCAAATCGAAGCGCGAGTCTCGACTTTTGGCCAAATTCCAGCAACACTCAAAACGGGAAAAGCGTCTATGGAGGATTGGGA
ATTGAGCGAGAGACGCTTCAGTCGGAGCTGTGAGCAAAATCTGGAATCGCAGGAGGACACATTTGGCAACACGAGGATGACGGCAAACTTCGGCAGCGCTTGTGTGCATTCGAGAT
TGCATTACTTCAGTTTTAAATACGCGACAGCGCGGAAAATAAGCAACAAATTCAGGGGTAAACGTGGCTGACCTTTCGGCCACGACATGACATACCATACCATAAC
AGCGCGAGCAGGAGCAGCAACATCACAAAGCAGCAGCACATCAGGCCACCAACGCGCAGCCACTGCAACTGCAACATTAACAGGCCACATTTCACGACGACGAGCAGC
ACTACGACAGCGCGGACGAGAGCTTGGCTATAACATTCGCAATTTGGTGATGCGCTGTTGACTTCGTATCATTAACAGCCACATCGTCAGATGCCCTCTCTCTCGCCCT
TTTCACTCTACTTGTGCTCGGATTCCAGCTTTGAGCTCTCTCGACAGTCGGCCCAAATAAAGCGCCAAATGCGCATGACATTCGGGTGGATAACACGGCGAGCTGGGGA
GAGCTGGCTGGATCTATCGCTGCTGCTCTCAAGGATTCATCTTCTGTCGAATATCTCTGGCCGCTGTCCTCGGCACATGCATTGGTCATCATTTAGTCGACGCCAAATCGA
AAGCTCGGGGTTGATAACCAATTAATCTTGTGTGTCGCTGGCGATGGCCGACATGCTGGTGCCCTCTGTGCGATGACATTAACGCCCTCGTGGAACTGTCGGCGGAAAGT
GAGTCTCGGACCGTCTATGTGCAACGCTGTACACGCGCTGGATGTTTACTTTTCCACGGCAGCATATTTGCACCTGTGCTGCATATCAGTGGACAGATACTACGCCATGTG
CGTCCACTGGAGTATCCATTGAATATGACACACAAAACGGTCTGCTTCTCATGCTGCCAAATGTTGGATCTCTCGGCCCTCATATCTTTCACGCCCATCTTTCTGGGCTGG
TACACGCGGAGGAGCAGCTCGCGGAGATTTCCCTGCATCGGCACAGTGCTCTGTTTGTGTCAACAGGCCCTACGCCCTCATCTCCAGTTCCGGTAGCTTCTGGTATACCGG
GCATTTGTGATCGCTGGTGTATGTAATCGGCCATCTTTAAGGAGGCGATTCTGTCAACGCAAGGCCCTCAGCCGACACAGCTCCAACATCTCTCTTAACAGCGTCCACATGGGCCA
CACCAGCGAGCCACCGACCGCTGAGCTATTCGATCATGACTGACTGTATCTGAAATGCCAATCTCGCCCGGGGAGGACGACGACAGTCGCGTTAGTAATCTGGAGCGGCCCGCG
GGCATCACCTTTCGCCGCGTGCACAGTCCGCTGCCAACTGTCCGCTCTGCGCGCCGACATCAGCAGCACACCGGCACACCGCAACCGCAACGCCAACGCCAACCGCG
ACAGCAACAATCGACCCACAGATTAACCACTAGTAGTAACCGTAGTATTACAGAGCAAACTCCAGATCTTGGCCAGAGGCCCGCCCTCCTCGTCCAGTTCCACTCGATTCTGGCA
CAACGCAAGACGGCGGCGGTACAGACTGTGGCAGCAGAGCAAGAACGGGAAGCGCGCTTCAAAACCCGATGACAGTACATCTGGTGGAACCCGCGGCTGTCGTCGACGCGCG
AAAGCATGGAAGGGCGGAACACAGGCGCCCGACCTTGGGCATCATCATGGGCTCTTTTGTGCTGCTGGCTGCCCTTCTTTCTGTGGTATGTATCATCATCTGCTCTGCG
GTCCGGCTCGCCATGTCCCGATGTGCTCGTGGTGGTGTATTCTGGATCGGTTACTTCACTCCACGCTAAATCCGCTTATATAGCCCTACTTTAATCGCGATTTTCGCGA
GGCATTCGCAATACGCTGGAGTGTGTCTGCCCTGTCTGGAGAAACGAAATCCGTACACGCGCTACTACTAGCTCTAG
(SEQ ID NO: 188)

Start ATG: 281 (Reverse strand: CAT)

MSGVNVADLLATTMTLPTTAAGAASATSAATSATNASHLQPATITGHSITTAARKRTTTPSSLPITSQFVDAISLTLSLTATSSDASYSPPSSSYLSLSDSTFEELLSTVGP
NTANGSDIADNDNQAELESWLDSLKGLFSSIIILAVLGNALVLIISQVRNKLRAVITNYFVUSLANVALCALMVALCANFNASVELSGGKMGFGFPMCNVNSLSDVYES
TASILLHCCISVDRYATVRLEYPLEPMTHTKVCFFILANVLPALISFTPIFLGWYTTEEHLREISLHPDQCSFVVUNKAYALISSVSFWIPGIVMLVMYWRIFKEAIRQR
KALSRSSNNILLNSVHMGHTQOPTSLSYLHPSDCDLNATSAAREETHSALSNIERPRGITFAAVPSPLPKCPLCGADISSTTGTTANATATANADSTIDTTVTSSKRSIHEQ
TPDLGQRPASSSSSTREFWHKRTAAITACMGQSKNRKRREFTGCSHCAGTGSVRPAKGNAEHKAATVIMQGVFLCWLPPFFLWYVITLSCGPACPCPDVLVVVLFWIGY
ENSTLNLPIIYAFNFDREAFRNTLECVLPCKLEKRNPNYAYYV*
(SEQ ID NO: 189)

Name: Beta Adrenergic/ Octopamine Receptor-like
Classification: G protein linked receptor

GTTCCTGGCTTACTTGGCTCTTGACTCTTTACCCAGAAATATGATTCTTGGCAGCAGGAGAAACAGTACGACTTGTTTTGGTCGTATTATGAGTGAATATTACCACATCATGCT
ATTTTGAACGCTTTTAAGAGCTCACACCCTTATGGACCCATTATAGAATATGCGGTGGCAAAATTTCCCAATTTTATCGAGGCTGGAAAAACTTTCCACACATGCCAATCAT
GTTTCGTTTCACTTATCATGGCTAAAACCTAGCAAAATGAGAAAAGTTTTTTTTATCCACAACATATCTGTATTAGTGGCCAGCAATCTGTATTGCTTTAGCTTAAGATGT
TTACTATTAAATAGAGCCAAAGTGGCAAAATGACATACGGCTTACAGAGAAAAATTAATTTCCAATTCAATTACATTTACGTACATATATCTTCTCTACAAGCAGACAGGATATT
GATCTAARATTAATAATAATAATAATAATTTAAATTAATAATAATTAATAATGATAGTAACAGAAATATATTTTGTATGAAGCCCTCATAGGTTATTGTGTGACTTT
TTTTCTCTCGCATCAAGTGAAGATGTTTACTATAATAGAGCCAGAGGCAAGAAAAAGTGGCAGCAATCAAGTGCTTTTATTTCGGGATCGCATGTGTAAGGCGCATTTCC
TGATCGAAGACTTTGGATACCTGATGCCATCCGCTGTAATGTGACGTCAGCCTGGCTGGGGATTCAATTTAAATTCATCTTGAAGTCTAATGCTTTAATGTTGTTGCTT
CACAGGCGAGTGGCCAATTTTGGGCGCTGCTGCTACGTTCTTCCGCTATGTGCCACAGCGCCGCAAGGATACGACAGCAATTAATAACGCCATTGATACAGTGGGG
AGGCCGATTTGCTTTTGTTTTCTTTTCCCAACTTGGGCGTGGAAAGGAGTGTGTGGCTGGCGTGGAGTGGATAGAGTCCACAATTAATCGACACAGCTAGTATGTGCA
CGGCGTCGCCATCACCATAGCCAACGACAGCAACGATGATGGCATCAATCAATCATTCATGGCTCATGTGTCGCCAGTCCCAATCAGAGTCCCTCCATTTGAAGTGGCGCATC
GGTATCGCCTCATCCCAATGGCCAAATCCCGCAGAGGCCCCGAGATGTGCTGCTGAAGAACGACAAATTTCAACACATGTGCGCCACTCTGCTGAACATAACGACGCCGAGA
ACTTTTCAAAATCTCTGGGTTCCAGCAAGCGGCAAAATGCGAGCAATGGCAGGATGTTCCGCCGTGCAAGATCCCTGACCTCGCAACGCCCTACCGGTTTGTATGT
ACTCATTTTTGTGGCCGGCGTGTGGGTAATCTTATCACCTGCATTTGTTATATCGCAAAATAACTTTATGCACAGCGCCACCAATTTCTATTTGTTCAACTGGCGGTTCCG
GACTTAATACTGTTGCTTACGAGTGAGATGGCATTTTGAAGGACACCATTCGCTATGCGTAAATGCGGCAAAATATCTGACGGCATCCCCAGGAGCTGTACAACCTTTGGT
ATCCGTAATCTGTATCTCTCAGGACGCCATGTGCATCGGCAGTGTGCTCGGAATGGCCGCAATGCAACGGTTCACCATCAGACGATCTACTGTGGAGCGATA
TATAGCAATCTGTCATCCATTCGGTGAGTTTTAAACATTTAAACGTTATATATATTTTAAATATTATTACATTATAAACAAATGTATAATTTCTTCAGTGTTATGCCC
TCGCTCTCATTAACAAAGTGGAATCATGATAATTAGGACAGAGAATATGAATAGATGAATTAACCTTGATAGCGCTGACACCGAAACATATGATATGAAGAACACAAAGTA
CCTCAAAATATAGGCGCAATAATTTATCAATCAACAGCTGTGGCTGTGGAATGTAAACAAAGTTAGGCTGCAGTCCGAATTTCCATCGTTTTAAATATACCAAT
TCATTAGCTTTTCAATTATAGACACATAATTAAGTCAGGTCCATAAATGAACCAAGTTCATACAGACATATGTAATTAAGTTCCAATAAATGCGTGTGAAGATAGAAGTTA
GTAGTACATTAATTTGCTTAGTACCAACATATTTGCAATTTTGGAAATGTTACACAAAACCTTTTCCCTATACAGGTTATCACCAGTCTCTTCCAATGTCTTGACCAATTTTG
GGTCAAGAATAACACAGCTCTCTTAAGCGAAAGGCTCTGGTTTTTCAATTCAGATGTGCATTTTCTTTAGTTCAGAACTCTTGATAGCTAGAGAAGTAAGAACTTAA
ACCTTACATTTGTGTTTTCAACACAAATTTCCGTTTCTTAGCATTTTTCACAAATTAATTTGTTACGAATACAATCACTTGGAGATGTGCTTGAAGATATGCAGACACAAA
GGGTGGTTTTCTATAAGAAAGGATTCCAGTCCGATTAACAAGTTAAGGCAAAAACCTTTTCATGCTCTGAAGTTGGGATTAATTTGTTGCGAAAAGAGTAAAGGTTT
CTATGTTTGGCAATTAATAGTATATAGTATTTTGAACAGAAAGGATATTCGGCATGTGCTAGCATGAAAAAGGCTGAAACCCATTCGACATCTTACCAAAAGAG
TTAAAGTATGATGCTATTAAATTTGGTATGAAACGTGTTGAATTCCTTCCCATTTACACCCACATTTCAAAGTAAATATTATGTTGCACATCAATCCCAAACCTTAAAGAG
AAAGTTAAAGCGTAAAGAGTGAGTGGTCACTGGCTGTACTTTTACAGACCGTAATATTTTCCATCTTATGCTGCGCCACCTTTATTCGAAACATCGACATCATCTTGCTT
TTAATCAATTAACATATAATGATGAGCTATGCTTTTCGTCATTTGTAGGACGACACCATGTCAAATTTGTGCGAGCGCTAAATTTATTTATGTCATTTGGCTGGCGGCTCT
CCTTTTGGCCCTTCCGCGAGGCATGCAATTTTCGGTGGTCTACCAGAACGAGGATACCTGTGCAAGGTGGGTGGGATGAATGTGAAAACCAACTGACGCATCATCTCGAT
GTCCTTTTCGCTGTTTATTCCTTATAGATGAAAGACGATTTTACGCCATGTGTTCCGCGTTTTCGGGATTTATTTTCTTCGGTGGACCTATGACGGCGATTGTGTACTGTA
CGTCTGATCGGAGTGAACCTGAAGAGAGCTCGACTGCTGCAATCGCTGCGCGAAGGACTTCGATGCGAATCGCGGCTAAATGCCAGGACGAGTACAGAATGTTG
GGTAGGTGAGTTAAGTTGGCGGCTTGATTGATTAAGACAGCAATATGCCCCACCGCCAAACAGAAACACAGACATCCAGCGGTGTCAAATTTTTCGCGCGAAAGT
TTAAATTTGAAATTTACTGTCAACGTTATGGCTGTGGCGAATCAACAGGGAATTCGTGATAGAGCCCGCCGCTATTGCTTCCGCCCCTTCAATTTTGGCTTGGCTCATGT
GGGCTGAACGCTGCTTCTGTTTTTTCCTCTTTTCCCTTCAACGCTGTGCTGAATTTTTCGAAACATCGGTGCTGATGTGTAAGCTCTTAAATGGCCATCTAGCCTATGAA
GCAAGGTGAAGCCAGGTGTCACTTCAAAAGTAATCAAGCCGGGACCTTAATGAATCAAGACTGCTGGGTGCACACAGCAAGCTTAATTTTGAAGAAAGTATTTTCTT
GAATCTATAGACATATGTCGTTAATGTCGCCAGTTTAAGCTGTAAATTTTAAGGAATTCAGACACATCTCGTGATTTATTCATATTTTATATATATGAA
TCTTATTAACACATACAGCAATAAATTTAAGTGTAAATCAGGCGCCAAATTTAAGATCCCAATAGCCGTTTTTCTTACAGCCTACGAAACAGCTTTAATTTACTGTG
CTTTAACTGAACACCCGACCATTGATTTTCAAAATTTACCATTTGGCCCTGGCTGCCAAATGGAATTTGATGACCAATTCAGATTAAGCCAGAGCCTTAACCAATAAT
ATTGGCGCCAGAGTGGAAAAAACAACCCGCTTCCAGGCTTAGTTATCATAGATGTTGGGATATTTATTCGTTTGTCAAGGAGTAATACCTTGGCTTTATCA
AAATGTGATATTTTACCAAGGCGGTTTGGCAACAACTTCGATTTGGGGGATATAAAGAAATAAGCACTTTTTCAGTGTGAGAGAGCCCTTTTCGGTACCTAGTTG

[illegible]

FIGURE SHEET 88

89/89

ATAGTAGAAAGTTTCCCTAAAGATCGATGTATAACTATTAATACATATTTTACTTAATCTAGTGTGCTTTTAAATGACAGTTTTTGTGTGCTTAAATTTTTACAAAATA
AATCTTAAATCCTTGCAAAGAGTTTCTTTTATGCAATGGAACATTTGTTTCATCGCTTGTAAATATCATTTACTTTATTATTATTATTCTAAACAGCTGCCAAAGCA
CGGAATATCTCAGGATCAGATATGATTGTCTATGATTTTGTACACTTCCATGTTTGTGATAACGCAAAAAGCACACACACCGTGATTGGCAATAAATGATAAATT
CTTTCCGCTCTGGTTTATACACTCACCGCATCGTCCATTCCCCCGTCCACTTTCCATCCGAGGGGCGACTCGCAGCGCAGCTCCATCCATGCTAACGGCGGTGACGAGG
TGGACGGCAGAGGTGGCAGTCCCGCCAGATGCGGGCCTTCCGCCAGCAGAGCTACTACCGCAGCTCCTCCAACGGCACAGCCGGACCGGTGCAGCTCCCTTAAAGGAGCA
GGTTGGCCTGCTGCACGTGGGTCCCGCAATGGGACGCCCGGTGGCTCCGTCTTAGCGCGGCCACGCCGAGTTGATCCGTAAAGGATCGGCTCTGTTGGCCCCGACAACCC
AGCTGTCTGAGGGAGCAGGAGCAGCAGCGTTTGTGCTGCTGCACGAGAAGCCCTCGACCCCTGGTGCTCAGCTACGACAGCCAGCGGGCGGAGTGGGCGTGGGCGTGGCCA
CGGCTCTGCTGGACAACAACGAGCGAGTGTGAGCGTGTGAGAGCAGCTGGCGATGGCGGTAGCGGCGGCGCGGTGGCGGTGGATGCACCGGACACAGTAGAAGCAGCAGC
AGCAACAGCATCAGCCGATGCGGATGCGAATGTGGCGATGGAGGAGCCCTGCTCCTGCTGCCATTGCTGAGCAGGGAGCTGGTGGCGGGCAGTTGGAGCTCCAAGTGCTT
CCAGTGCAGTGAGGCGACTGCCAGTGGATTTGATCCGGCTGGCTGAGTGGCCGGAGCTGGGGATTCCAGCGATTCCCTCGGTCAACGTGGGGAGTGGCGTGCATGATTAGAT
CACCGGGCGTGCACATGTGCTTAGATAGTGTAACTCGAGAGTTAAGTCTTCAAACACAATGGGTGTGCTTAGAAGTAGCGTAGGATTAGAGCTAGTCCGTACGAGTGGT
TGCCCTACACTGGGAGAACTCGTGGTCACTTTGCATGTTCTCTGAATGTTCTACTAATCTTGAATCAACAAGGATACAAATATAAGAGATAATATCTATGTACATCCCA
TAAAAACATAAAAAATGCAATTAAATCTGGTTGTAAAAAAATAGATTTAACTTGAATGTTTCAAATGTTGCCAATTCGCCAGCGTGAATCCGATATAAGCAACAATA
ATTACAAATTATAATCAATGTGCAATCAGAGTCAGTTACAATTTACAATTATATGACTGAATAATTGGACTATGAATATATACGACTTCAAATGAGTTTCTCTCATTAGCAGG
GCAACTTTGTAATGTTAATTAACAGCGTTTTTCAGTTTTTCATTTTTGTTCCCCAGCGAGTAACCAAAATCGATTCTATAATTGTGTAATAAGAAATCATCTAAAAAC
AGGGTAAATCTAACTATTGTAAATACATTTACTCATTATTTGAGAAGAATATTGTAAAAAAAAGATTGTAAGATTAAATAAATATAGTTTCATGTTCAACGACACAA
GGAATTATTTGATTATTTAAAGTACTAAAAA
(SEQ ID NO: 196)

Exon: 1001..1210
Exon: 1307..1774
Exon: 6357..6540
Exon: 6605..6810
Exon: 7683..7834
Exon: 8466..8889
Start ATG: 1001

Transcript No. : CT29989

ATGGAGGACGAGTGGGGCTCCTTTGATCGTCTGCCAGTGTTCGAGTGCCTCGATGGATTGGAGACGGAAAACGAGGTGGTCAGCAATTGGTCCACACTGGCCAACTTCA
CGCGACTTGTGGCTGGTGCCGCTCCTGAAATCGTCAACTATACGCTCAACATGATCGAGCTGGGTGTGGGCATGGCCACGGATATATCCAATTTGAGCGTTCCGGAGCACGT
GATGGATCAGCACCTCAACTATCCCGATCCGGTTGCTGAAAGTGATGTGCTGGCGGTAATGGCACTGTTCTCTCTGCTGGGCAACCTGGTGACCATCTGGAATATCTAC
AAAACCGCATCTCAAGAAAGAACTCACGCGACACGTGGAGTGTCTACTCACTGATGTTCCATCTGTCCATCGCCGATGCTCTGGTCACTGGTTCTGCATTATCGGGG
AGGCCGCGTGGTGTACACGCTCCAGTGGCTGGCCAAATGAGCTCACCTGCAAGCTGGTGAAGCTCTTCCAGATGTTAGCCTCTACCTGAGCACCTATGTCCTGGTCTCAT
CGGAGTGGACCGCTGGATAGCGGTCAAGTATCCGATGAAGTCGCTCAACATGGGCAAGAGGTGTCATAGGCTACTTGGCGGTACTTACATCTCTGCTGGTGGTCTAGCTTG
CCACAGTCTCTCATCTTCCATGTAGCGCGTGGCCCATTCGTGGAGGAGTTTACCAGTGCCTCACCCACGGATTCTACACGGCGGATTGGCAGGAGCAGATGTACGCCACCT
TCACGCTGGTCTTCACTTCTCTGCTGCGCTGTGCATCTGTTTGGCACCTACATGTCCACCTTCCGCACCATTTCCAGCAGCGAAAAGATGTTCAGGGATCAAAGTTGGC
CAACTACTCAACGGCCAAATGGCCACACAGACGAATCGCCAGAGGCTGATACACAAGGCCAAGATGAAGTCGCTTCGCATATCCGTGGTGATCATCATAGCGTTTCTCATC
TGCTGGACGCCCTACTACGTCATGATGATTATGTTTCTTCAATCCGGACAAAAGGCTGGGCGACGATCTGCAGGACGCCATCTTCTTCTTCGGCATGTCAACAGCC
TGGTCAACCCACTCATCTACGGTGCCTTCCACCTGTGCTCTGGCAAGGGGGCAAGTCGAGCGGCGGGGCGGCAACACACGCTACAGCTTGAACAGGGGCGACTCGCA
GCGCACTCCATCCATGCTAACGGCGGTGACGCAAGTGGACGGGACAGGTGGCAGTTCCCGCCAGATGCGGGCCTTCCGCCAGCAGAGCTACTACCGCAGCTCTTCAACGGC
ACAGCCGACCGGGTGCAGCTCCTTTAAGGAGCAGGTGGCCCTGCTGCACGTGGGTCCCGGCAATGGGACGCCCGGTGGCTCCGTCTTACGGCGGCCACGCCGAGTGA
TCCGTAAAGGATCGGCTCTGTTGGCCGACAAACCCAGCTGTCTGAGGGAGCAGGAGCACCAGCAGCGTTTGTGCTGCACGAGAAGCCCTCGACCTGGTGTGCTGACGACG
CAGCCAGCGGGCGGAGTGGCGTGGCGTGGCCAGCGGTCTGCTGGACAACACAGAGCGAGTGTGAGCGGTGTA
(SEQ ID NO: 197)

Start ATG: 1

MEDEWGSFDRLPSPVPSASMDLETENEVVSNWSTLANFTRLVAGAAPEIVNYTNLMIDVGVGMATDISNLSVPEHVMOHAPQLSRSGLLKVYVLAVMALFSLGNLLTIWNIY
KTRISRRNSRHTWSAIYSLMFHLSIADVLVTFWCIIGEAAWCYTVQWLANELTCKLVKLFQMFSLYLYLVLVIGVDRWIAVKYPMKSLNMAKCHRLLGGTYILSLVLSL
PQFFIFHVARGPFVEEFYQCVTHGFYADWQEQMYATFTLVFTFLLPLCLFPGTYMSTFRTISSSEKMFQGSKLANYSTAKLPTQNRQRLIHKAKMSLRISVVIITAFLI
CWTPTYVMMIMFNLNPDKRLGDDLDQDAIEFFGMSNSLVNPLIYGAHFHLCFGKGGKSSGGGNNNAYS LNREGDSQRTPSMLTAVTQVDGTGSSSQMRFRQOSYRSSNG
TAGPGAAPFKEQVGLLHVGPNGTPGGSVSSGATPQLIRKGSALLARQPSCLREQEHQRLLLHEKPSTLVLSYDSQRGGVGVGASGLDNNERVSSV*
(SEQ ID NO: 198)

Name: GONADOTROPIN-RELEASING HORMONE RECEPTOR-like
Classification: G_protein_linked_receptor

INTERNATIONAL SEARCH REPORT

 Application No
 PCT/US 01/09341

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C12N15/12 C07K14/435 C07K14/705 C07K16/18 C12N5/10 C12Q1/68 G01N33/50		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 7 C07K G01N C12N C12Q		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the International search (name of data base and, where practical, search terms used) SEQUENCE SEARCH, EPO-Internal, BIOSIS, WPI Data, PAJ		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE EMBL SEQUENCE LIBRARY [Online] 6 January 2000 (2000-01-06) ADAMS, M. AND VENTER, J.C.: "Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in ordered pieces" XP002182628 accession no AC020076 ---	1,2,4,5
A	WO 94 08006 A (ZYMOGENETICS INC) 14 April 1994 (1994-04-14) the whole document --- -/--	
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "8" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
13 November 2001		15. 02. 2002
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer Holtorf, S

INTERNATIONAL SEARCH REPORT

Application No

PCT/US 01/09341

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>LI X-J ET AL: "CLONING, FUNCTIONAL EXPRESSION, AND DEVELOPMENTAL REGULATION OF A NEUROPEPTIDE Y RECEPTOR FROM DROSOPHILA MELANOGASTER" JOURNAL OF BIOLOGICAL CHEMISTRY, AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD, US, vol. 267, no. 1, 5 January 1992 (1992-01-05), pages 9-12, XP000877443 ISSN: 0021-9258 the whole document</p> <p>---</p>	
A	<p>WO 99 01468 A (DEN HEUVEL MARCEL VAN ;INGHAM PHILIP W (GB); ONTOGENY INC (US)) 14 January 1999 (1999-01-14) the whole document</p> <p>---</p>	
A	<p>HAUSER FRANK ET AL: "Molecular cloning, genomic organization and developmental regulation of a novel receptor from Drosophila melanogaster structurally related to gonadotropin-releasing hormone receptors from vertebrates." BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, vol. 249, no. 3, 28 August 1998 (1998-08-28), pages 822-828, XP002182627 ISSN: 0006-291X the whole document</p> <p>---</p>	
A	<p>FENG G ET AL: "CLONING AND FUNCTIONAL CHARACTERIZATION OF A NOVEL DOPAMINE RECEPTOR FROM DROSOPHILA MELANOGASTER" JOURNAL OF NEUROSCIENCE, NEW YORK, NY, US, vol. 15, no. 12, 15 June 1995 (1995-06-15), pages 3925-3933, XP002919142 ISSN: 0270-6474 the whole document</p> <p>---</p>	
P,X	<p>ADAMS M D ET AL: "THE GENOME SEQUENCE OF DROSOPHILA MELANOGASTER" SCIENCE, AMERICAN ASSOCIATION FOR THE ADVANCEMENT OF SCIENCE,, US, vol. 287, no. 5461, 24 March 2000 (2000-03-24), pages 2185-2195, XP000961051 ISSN: 0036-8075 the whole document</p> <p>---</p>	1,2,4,5
	<p>---</p> <p>-/--</p>	

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 01/09341

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-20 partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

Invention 1: claims 1-20 partially

Isolated protein consisting or comprising an amino acid sequence as characterized by SEQID3, or an allelic variant or an ortholog of said amino acid sequence wherein said variant or ortholog is encoded by a nucleic acid molecule that hybridizes to the nucleic acid molecule as characterized by SEQID1 or 2; an antibody that binds to said protein; furthermore a nucleic acid molecule consisting or comprising of a nucleotide sequence that

- 1) encodes the amino acid sequence of SEQID3
- 2) encodes an allelic variant or an ortholog of an amino acid sequence of SEQID3 wherein said nucleotide sequence hybridizes to SEQID 1 or 2
- 3) encodes a fragment of said SEQID3
- 4) is the complement of the nucleotides of 1) to 3)

The recombinant expression of the same in host cells and methods for the detection of said proteins or said nucleic acids in a sample with the help of an agent that binds to said protein or an oligonucleotide and kits that contain such agent or oligonucleotide.

Furthermore, a method to identify an agent that binds to said protein by detecting a complex formed by an agent and the said protein.

Invention 2-66: claims 1-20 partially

as invention one but referring to the protein and nucleic acid sequences as characterized by SEQIDs 6,9,....,192,195,198; SEQIDs 4,7,....,190,193,196 and SEQIDs 5,8,....,191,194,197, respectively.

INTERNATIONAL SEARCH REPORT

Application No
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C. (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
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